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OM protein - protein search, using sw model

Run on: March 28, 2006, 11:16:29 ; Search time 29.5986 Seconds
(without alignments)
1661.969 Million cell updates/sec

Title: US-10-659-000-4
Perfect score: 3231
Sequence: 1 STIEEQAKTFLDKFNHEARD.....WLKDNQKNSFVGNSTDMSPY 595

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 572060 seqs, 82675679 residues

Total number of hits satisfying chosen parameters: 572060

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA:*
1: /cgn2_6/prodata/1/iaa/5 COMB.pep.*
2: /cgn2_6/prodata/1/iaa/6 COMB.pep.*
3: /cgn2_6/prodata/1/iaa/H COMB.pep.*
4: /cgn2_6/prodata/1/iaa/PCTUS COMB.pep.*
5: /cgn2_6/prodata/1/iaa/RE COMB.pep.*
6: /cgn2_6/prodata/1/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	3231	100.0	805	2	US-08-989-299-2
2	3231	100.0	805	2	US-10-158-847-142
3	3231	100.0	805	2	US-09-407-427-2
4	3231	100.0	805	2	US-09-635-501-2
5	3231	100.0	805	2	US-09-635-501-106
6	3231	100.0	805	2	US-10-158-825-142
7	3231	100.0	819	2	US-09-949-016-11284
8	2987	92.4	681	2	US-10-158-847-140
9	2987	92.4	681	2	US-10-158-825-140
10	2987	92.4	711	2	US-10-158-847-138
11	2987	92.4	711	2	US-10-158-825-138
12	1335	41.3	732	1	US-08-481-626-2
13	1335	41.3	732	2	US-08-989-299-4
14	1335	41.3	732	2	US-09-407-427-4
15	1335	41.3	732	2	US-09-635-501-4
16	1335	41.3	1265	2	US-09-964-899-19
17	1335	41.3	1306	2	US-08-989-299-7
18	1335	41.3	1306	2	US-09-407-427-7
19	1335	41.3	1306	2	US-09-635-501-7
20	1334	41.3	732	2	US-08-989-299-5
21	1334	41.3	732	2	US-09-407-427-5
22	1334	41.3	732	2	US-09-635-501-5
23	1334	41.3	1312	2	US-08-989-299-8
24	1334	41.3	1312	2	US-09-407-427-8
25	1334	41.3	1312	2	US-09-635-501-8
26	1310	40.5	1313	2	US-08-989-299-9
27	1310	40.5	1313	2	US-09-407-427-9

28	1310	40.5	1313	2	US-09-635-501-9	Sequence 9, Appli
29	1281	39.6	737	2	US-08-989-299-6	Sequence 6, Appli
30	1281	39.6	737	2	US-09-407-427-6	Sequence 6, Appli
31	1281	39.6	737	2	US-09-635-501-6	Sequence 6, Appli
32	1281	39.6	1310	2	US-08-989-299-10	Sequence 10, Appli
33	1281	39.6	1310	2	US-09-407-427-10	Sequence 10, Appli
34	1281	39.6	1310	2	US-09-635-501-10	Sequence 10, Appli
35	1073	33.2	615	2	US-08-989-299-11	Sequence 11, Appli
36	1073	33.2	615	2	US-09-407-427-11	Sequence 11, Appli
37	1073	33.2	615	2	US-09-635-501-11	Sequence 11, Appli
38	989	30.6	694	2	US-09-440-325A-1	Sequence 1, Appli
39	989	30.6	694	2	US-09-846-996A-1	Sequence 1, Appli
40	989	30.6	694	2	US-10-246-085A-1	Sequence 1, Appli
41	981	30.4	590	2	US-09-902-540-10939	Sequence 10939, A
42	677	21.0	149	2	US-09-621-976-3897	Sequence 3897, Ap
43	635.5	19.7	907	2	US-08-989-299-12	Sequence 12, Appli
44	635.5	19.7	907	2	US-09-407-427-12	Sequence 12, Appli
45	635.5	19.7	907	2	US-09-635-501-12	Sequence 12, Appli

ALIGNMENTS

RESULT 1
US-08-989-299-2
; Sequence 2, Application US/08989299
; Patent No. 6194556
; GENERAL INFORMATION:
; APPLICANT: Acton, Susan L.
; APPLICANT: Robinson, Keith E.
; TITLE OF INVENTION: ANGIOTENSIN CONVERTING ENZYME HOMOLOG
; TITLE OF INVENTION: AND THERAPEUTIC AND DIAGNOSTIC USES THEREFOR
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: FOLEY, HOAG & ELIOT LLP
; STREET: One Post Office Square
; CITY: Boston
; STATE: MA
; COUNTRY: USA
; ZIP: 02109-2170
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/989,299
; FILING DATE: 11-DEC-1997
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Arnold E., Beth
; REGISTRATION NUMBER: 35,430
; REFERENCE/DOCKET NUMBER: MIA-025.01
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-832-1000
; TELEFAX: 617-832-7000
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 805 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-989-299-2

Query Match 100.0%; Score 3231; DB 2; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.8e-309; Indels 0; Gaps 0;
Matches 595; Conservative 0; Mismatches 0;

Qy	1	STIEEQAKTFLDKFNHEARDLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLEQST 60
Db	19	STIEEQAKTFLDKFNHEARDLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLEQST 78
Qy	61	LAQMTPLOEIQNLTVKQLQALQQNGSSVLSEDKSKRLNTILNTWTSTYSTGKVCNPNP 120

Db 79 LAQMPLOBIQNLTVKQLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 138
Qy 121 QSCLLLEPGNETMANSNDYNERLWAWESRSEVGVKQLRPLVEEYVVLKNEWARANHVED 180
Db 139 QSCLLLEPGNETMANSNDYNERLWAWESRSEVGVKQLRPLVEEYVVLKNEWARANHVED 198
Qy 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVBHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
Db 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVBHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
Qy 241 IGCLPAHLGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRI FKEAEKFFVSV 300
Db 259 IGCLPAHLGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRI FKEAEKFFVSV 318
Qy 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVMTWDDFLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVMTWDDFLTAHHEMGH 378
Qy 361 IOYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKXIGLLSPDPFQEDNTEINF 420
Db 379 IOYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKXIGLLSPDPFQEDNTEINF 438
Qy 421 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIIVGVVPEVPHDETYC 480
Db 439 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIIVGVVPEVPHDETYC 498
Qy 481 DPASLPHVSNDSYFRIRYTRTYQFOFQALCOAAKHEGPHLKCDISNSTEAGQKLFNML 540
Db 499 DPASLPHVSNDSYFRIRYTRTYQFOFQALCOAAKHEGPHLKCDISNSTEAGQKLFNML 558
Qy 541 RLKSEPTWTLALENVVGAKNMVRPLLNYFEPFLFTWLKDQKNKSFVGMSTWSPY 595
Db 559 RLKSEPTWTLALENVVGAKNMVRPLLNYFEPFLFTWLKDQKNKSFVGMSTWSPY 613

RESULT 2
US-10-158-847-142
; Sequence 142, Application US/10158847
; Patent No. 6592865
; GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACE-2 Activity
; FILE REFERENCE: PF557
; CURRENT APPLICATION NUMBER: US/10/158,847
; PRIOR FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/295,004
; PRIOR FILING DATE: 2001-06-04
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 142
; LENGTH: 805
; TYPE: PRT
; ORGANISM: homo sapiens
US-10-158-847-142

Query Match 100.0%; Score 3231; DB 2; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.8e-309;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 STIEEQAKTFDKFNHEADLFYQSSLASWNTNTI TEENVQNMNAGDKWSAFLKEQST 60
Db 19 STIEEQAKTFDKFNHEADLFYQSSLASWNTNTI TEENVQNMNAGDKWSAFLKEQST 78
Qy 61 LAQMPLOBIQNLTVKQLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 120
Db 79 LAQMPLOBIQNLTVKQLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 138
Qy 121 QSCLLLEPGNETMANSNDYNERLWAWESRSEVGVKQLRPLVEEYVVLKNEWARANHVED 180
Db 139 QSCLLLEPGNETMANSNDYNERLWAWESRSEVGVKQLRPLVEEYVVLKNEWARANHVED 198
Qy 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVBHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 240

Db 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVBHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
Qy 241 IGCLPAHLGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRI FKEAEKFFVSV 300
Db 259 IGCLPAHLGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRI FKEAEKFFVSV 318
Qy 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVMTWDDFLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVMTWDDFLTAHHEMGH 378
Qy 361 IOYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKXIGLLSPDPFQEDNTEINF 420
Db 379 IOYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKXIGLLSPDPFQEDNTEINF 438
Qy 421 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIIVGVVPEVPHDETYC 480
Db 439 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIIVGVVPEVPHDETYC 498
Qy 481 DPASLPHVSNDSYFRIRYTRTYQFOFQALCOAAKHEGPHLKCDISNSTEAGQKLFNML 540
Db 499 DPASLPHVSNDSYFRIRYTRTYQFOFQALCOAAKHEGPHLKCDISNSTEAGQKLFNML 558
Qy 541 RLKSEPTWTLALENVVGAKNMVRPLLNYFEPFLFTWLKDQKNKSFVGMSTWSPY 595
Db 559 RLKSEPTWTLALENVVGAKNMVRPLLNYFEPFLFTWLKDQKNKSFVGMSTWSPY 613

RESULT 3
US-09-407-427-2
; Sequence 2, Application US/09407427
; Patent No. 6610497
; GENERAL INFORMATION:
; APPLICANT: Acton, Susan L.
; APPLICANT: Robison, Keith E.
; TITLE OF INVENTION: ANGIOTENSIN CONVERTING ENZYME HOMOLOG AND THERAPEUTIC
; FILE REFERENCE: MNI-132CP2
; CURRENT APPLICATION NUMBER: US/09/407,427
; CURRENT FILING DATE: 1999-09-29
; PRIOR APPLICATION NUMBER: 09/163,648
; PRIOR FILING DATE: 1998-09-30
; PRIOR APPLICATION NUMBER: 08/989,299
; PRIOR FILING DATE: 1997-12-11
; NUMBER OF SEQ ID NOS: 14
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 2
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-407-427-2

Query Match 100.0%; Score 3231; DB 2; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.8e-309;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 STIEEQAKTFDKFNHEADLFYQSSLASWNTNTI TEENVQNMNAGDKWSAFLKEQST 60
Db 19 STIEEQAKTFDKFNHEADLFYQSSLASWNTNTI TEENVQNMNAGDKWSAFLKEQST 78
Qy 61 LAQMPLOBIQNLTVKQLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 120
Db 79 LAQMPLOBIQNLTVKQLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 138
Qy 121 QSCLLLEPGNETMANSNDYNERLWAWESRSEVGVKQLRPLVEEYVVLKNEWARANHVED 180
Db 139 QSCLLLEPGNETMANSNDYNERLWAWESRSEVGVKQLRPLVEEYVVLKNEWARANHVED 198
Qy 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVBHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
Db 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVBHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
Qy 241 IGCLPAHLGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRI FKEAEKFFVSV 300

Db 259 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDAQWDAQRIFKAEKFFVSV 318
Qy 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMCH 360
Db 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMCH 378
Qy 361 IOYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 420
Db 379 IOYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 438
Qy 421 LLKQALTIIVGLTPTFTYMLKRWMMVFKGEIPKQDMKKWEMKREIVGVVBPVPHDETYC 480
Db 439 LLKQALTIIVGLTPTFTYMLKRWMMVFKGEIPKQDMKKWEMKREIVGVVBPVPHDETYC 498
Qy 481 DPASLFHVSNDYSFIRYRTLYQFOFQALCOAAKHEGPHLKCDISNSTEAGQKLFNNL 540
Db 499 DPASLFHVSNDYSFIRYRTLYQFOFQALCOAAKHEGPHLKCDISNSTEAGQKLFNNL 558
Qy 541 RLKSEPTWLALENVVGAKNNVRPLLNYFBLFTWLKDQNKNSFVGWSTDWSPY 595
Db 559 RLKSEPTWLALENVVGAKNNVRPLLNYFBLFTWLKDQNKNSFVGWSTDWSPY 613

RESULT 4

US-09-635-501-2
; Sequence 2, Application US/09635501
; Patent No. 6884771
; GENERAL INFORMATION:
; APPLICANT: Acton, Susan L. et al.
; TITLE OF INVENTION: ANGIOTENSIN CONVERTING ENZYME HOMOLOG AND THERAPEUTIC
; FILE REFERENCE: MNI-132CP3
; CURRENT FILING DATE: 2000-08-09
; PRIOR APPLICATION NUMBER: US/09/635,501
; PRIOR FILING DATE: 09/407,427
; PRIOR APPLICATION NUMBER: 09/163,648
; PRIOR FILING DATE: 1998-09-30
; PRIOR APPLICATION NUMBER: 08/989,299
; PRIOR FILING DATE: 1997-12-11
; NUMBER OF SEQ ID NOS: 107
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 2
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-635-501-2

Query Match 100.0%; Score 3231; DB 2; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.8e-309;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 STIEBOAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLEKQST 60
Db 19 STIEBOAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLEKQST 78
Qy 61 LAQMPLOEIQNLTKVQLQALQNGSSVLSSEKSKRLNTILNTWSTIYSTGKVCNPNP 120
Db 79 LAQMPLOEIQNLTKVQLQALQNGSSVLSSEKSKRLNTILNTWSTIYSTGKVCNPNP 138
Qy 121 QECLLLEPGLNEMANSLDYNERLWAWESRSEVKGQRLPLYEYVVLKNEARAHYED 180
Db 139 QECLLLEPGLNEMANSLDYNERLWAWESRSEVKGQRLPLYEYVVLKNEARAHYED 198
Qy 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEKPLYEHLHAYVRAKLMNAYSISYP 240
Db 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEKPLYEHLHAYVRAKLMNAYSISYP 258
Qy 241 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDAQWDAQRIFKAEKFFVSV 300
Db 259 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDAQWDAQRIFKAEKFFVSV 318

Qy 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMCH 360
Db 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMCH 378
Qy 361 IOYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 420
Db 379 IOYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 438
Qy 421 LLKQALTIIVGLTPTFTYMLKRWMMVFKGEIPKQDMKKWEMKREIVGVVBPVPHDETYC 480
Db 439 LLKQALTIIVGLTPTFTYMLKRWMMVFKGEIPKQDMKKWEMKREIVGVVBPVPHDETYC 498
Qy 481 DPASLFHVSNDYSFIRYRTLYQFOFQALCOAAKHEGPHLKCDISNSTEAGQKLFNNL 540
Db 499 DPASLFHVSNDYSFIRYRTLYQFOFQALCOAAKHEGPHLKCDISNSTEAGQKLFNNL 558
Qy 541 RLKSEPTWLALENVVGAKNNVRPLLNYFBLFTWLKDQNKNSFVGWSTDWSPY 595
Db 559 RLKSEPTWLALENVVGAKNNVRPLLNYFBLFTWLKDQNKNSFVGWSTDWSPY 613

RESULT 5

US-09-635-501-106
; Sequence 106, Application US/09635501
; Patent No. 6884771
; GENERAL INFORMATION:
; APPLICANT: Acton, Susan L. et al.
; TITLE OF INVENTION: ANGIOTENSIN CONVERTING ENZYME HOMOLOG AND THERAPEUTIC
; FILE REFERENCE: MNI-132CP3
; CURRENT FILING DATE: 2000-08-09
; PRIOR APPLICATION NUMBER: US/09/635,501
; PRIOR FILING DATE: 09/407,427
; PRIOR APPLICATION NUMBER: 09/163,648
; PRIOR FILING DATE: 1998-09-30
; PRIOR APPLICATION NUMBER: 08/989,299
; PRIOR FILING DATE: 1997-12-11
; NUMBER OF SEQ ID NOS: 107
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 106
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-635-501-106

Query Match 100.0%; Score 3231; DB 2; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.8e-309;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 STIEBOAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLEKQST 60
Db 19 STIEBOAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLEKQST 78
Qy 61 LAQMPLOEIQNLTKVQLQALQNGSSVLSSEKSKRLNTILNTWSTIYSTGKVCNPNP 120
Db 79 LAQMPLOEIQNLTKVQLQALQNGSSVLSSEKSKRLNTILNTWSTIYSTGKVCNPNP 138
Qy 121 QECLLLEPGLNEMANSLDYNERLWAWESRSEVKGQRLPLYEYVVLKNEARAHYED 180
Db 139 QECLLLEPGLNEMANSLDYNERLWAWESRSEVKGQRLPLYEYVVLKNEARAHYED 198
Qy 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEKPLYEHLHAYVRAKLMNAYSISYP 240
Db 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEKPLYEHLHAYVRAKLMNAYSISYP 258
Qy 241 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDAQWDAQRIFKAEKFFVSV 300
Db 259 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDAQWDAQRIFKAEKFFVSV 318
Qy 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMCH 360
Db 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMCH 378

QY 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGLLSPDFQEDNETINF 420
DB 379 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGLLSPDFQEDNETINF 438
QY 421 LLKQALTIIVGTLPTFTYMLEKRWMPFKGBIPKQDQWKKWEMKREIVGVVPEVPHDETYC 480
DB 439 LLKQALTIIVGTLPTFTYMLEKRWMPFKGBIPKQDQWKKWEMKREIVGVVPEVPHDETYC 498
QY 481 DPASLFHVSNDYSFIRYTRTYLQFQFOEALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 540
DB 499 DPASLFHVSNDYSFIRYTRTYLQFQFOEALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 558
QY 541 RLKSEPTWTLALENVVGAKNMVRPLLNYFEPFLFTWLKDQNKNSFVGWSTWSPY 595
DB 559 RLKSEPTWTLALENVVGAKNMVRPLLNYFEPFLFTWLKDQNKNSFVGWSTWSPY 613

RESULT 6
US-10-158-825-142
; Sequence 142, Application US/10158825
; Patent No. 6900033
; GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACS-2 Activity
; FILE REFERENCE: PF555
; CURRENT APPLICATION NUMBER: US/10/158,825
; CURRENT FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/294,976
; PRIOR FILING DATE: 2001-06-04
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 142
; LENGTH: 805
; TYPE: PRT
; ORGANISM: homo sapiens
US-10-158-825-142

Query Match 100.0%; Score 3231; DB 2; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.8e-309; Indels 0; Gaps 0;
Matches 595; Conservative 0; Mismatches 0;
QY 1 STIEEQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWSAFLEKEQST 60
DB 19 STIEEQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWSAFLEKEQST 78
QY 61 LAQMYPLQEIQLTVKQLQALQONGSSVLSBDSKRLNTILNTMTSTIYSTGKVCNPNP 120
DB 79 LAQMYPLQEIQLTVKQLQALQONGSSVLSBDSKRLNTILNTMTSTIYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEYVVLKNEMARANHYED 180
DB 139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEYVVLKNEMARANHYED 198
QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRACKLNAYPSYISP 240
DB 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRACKLNAYPSYISP 258
QY 241 IGCLPAHLGLDMWGRFNTNLSLTVPGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 300
DB 259 IGCLPAHLGLDMWGRFNTNLSLTVPGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGDPRILMCTKVTDWDDFLTAHHEMGH 360
DB 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGDPRILMCTKVTDWDDFLTAHHEMGH 378
QY 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGLLSPDFQEDNETINF 420
DB 379 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGLLSPDFQEDNETINF 438
QY 421 LLKQALTIIVGTLPTFTYMLEKRWMPFKGBIPKQDQWKKWEMKREIVGVVPEVPHDETYC 480
DB 439 LLKQALTIIVGTLPTFTYMLEKRWMPFKGBIPKQDQWKKWEMKREIVGVVPEVPHDETYC 498

QY 481 DPASLFHVSNDYSFIRYTRTYLQFQFOEALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 540
DB 499 DPASLFHVSNDYSFIRYTRTYLQFQFOEALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 558
QY 541 RLKSEPTWTLALENVVGAKNMVRPLLNYFEPFLFTWLKDQNKNSFVGWSTWSPY 595
DB 559 RLKSEPTWTLALENVVGAKNMVRPLLNYFEPFLFTWLKDQNKNSFVGWSTWSPY 613
RESULT 7
US-09-949-016-11284
; Sequence 11284, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11284
; LENGTH: 819
; TYPE: PRT
; ORGANISM: Human
US-09-949-016-11284

Query Match 100.0%; Score 3231; DB 2; Length 819;
Best Local Similarity 100.0%; Pred. No. 2.9e-309; Indels 0; Gaps 0;
Matches 595; Conservative 0; Mismatches 0;
QY 1 STIEEQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWSAFLEKEQST 60
DB 33 STIEEQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWSAFLEKEQST 92
QY 61 LAQMYPLQEIQLTVKQLQALQONGSSVLSBDSKRLNTILNTMTSTIYSTGKVCNPNP 120
DB 93 LAQMYPLQEIQLTVKQLQALQONGSSVLSBDSKRLNTILNTMTSTIYSTGKVCNPNP 152
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEYVVLKNEMARANHYED 180
DB 153 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEYVVLKNEMARANHYED 212
QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRACKLNAYPSYISP 240
DB 213 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRACKLNAYPSYISP 272
QY 241 IGCLPAHLGLDMWGRFNTNLSLTVPGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 300
DB 273 IGCLPAHLGLDMWGRFNTNLSLTVPGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 332
QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGDPRILMCTKVTDWDDFLTAHHEMGH 360
DB 333 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGDPRILMCTKVTDWDDFLTAHHEMGH 392
QY 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGLLSPDFQEDNETINF 420
DB 393 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGLLSPDFQEDNETINF 452
QY 421 LLKQALTIIVGTLPTFTYMLEKRWMPFKGBIPKQDQWKKWEMKREIVGVVPEVPHDETYC 480
DB 453 LLKQALTIIVGTLPTFTYMLEKRWMPFKGBIPKQDQWKKWEMKREIVGVVPEVPHDETYC 512
QY 481 DPASLFHVSNDYSFIRYTRTYLQFQFOEALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 540

Db 513 DPASLFHVSNDYSFIRYTRTLVQFQFQALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 572

Qy 541 RLKGSBPWTALLENVVGAKNMVRLPLNYFEBLFTWLKDQNKNSFVGNSTDWSPY 595

Db 573 RLKGSBPWTALLENVVGAKNMVRLPLNYFEBLFTWLKDQNKNSFVGNSTDWSPY 627

RESULT 8

US-10-158-847-140

; Sequence 140, Application US/10158847

; Patent No. 6592865

; GENERAL INFORMATION:

; APPLICANT: Tom Parry et al.

; TITLE OF INVENTION: Method and Compositions for Modulating ACE-2 Activity

; FILE REFERENCE: PF557

; CURRENT APPLICATION NUMBER: US/10/158,847

; PRIOR FILING DATE: 2002-06-03

; PRIOR APPLICATION NUMBER: 60/295,004

; PRIOR FILING DATE: 2001-06-04

; NUMBER OF SEQ ID NOS: 158

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 140

; LENGTH: 681

; TYPE: PRT

; ORGANISM: homo sapiens

; FEATURE:

; NAME/KEY: MISC FEATURE

; LOCATION: (219)..(219)

; OTHER INFORMATION: Xaa equals any amino acid

; FEATURE:

; NAME/KEY: MISC FEATURE

; LOCATION: (240)..(240)

; OTHER INFORMATION: Xaa equals any amino acid

; FEATURE:

; NAME/KEY: MISC FEATURE

; LOCATION: (499)..(499)

; OTHER INFORMATION: Xaa equals any amino acid

US-10-158-847-140

Query Match 92.4%; Score 2987; DB 2; Length 681;

Best Local Similarity 99.3%; Pred. No. 2.5e-285;

Matches 548; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 44 MNNAGDKWSAFLKEQSTLAQMYPLQEIQLTVKQLQALQOQSSVLSDEKSKRLNTILN 103

Db 1 MNNAGDKWSAFLKEQSTLAQMYPLQEIQLTVKQLQALQOQSSVLSDEKSKRLNTILN 60

Qy 104 TMSITYSTGKVCNPNPQECILLPEGLNEIMANSLDYNERLWAWESWRSVEVGKQLRPLYE 163

Db 61 TMSITYSTGKVCNPNPQECILLPEGLNEIMANSLDYNERLWAWESWRSVEVGKQLRPLYE 120

Qy 164 EYVLKNEMARANHYEDYDGYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEIKPLYEHLH 223

Db 121 EYVLKNEMARANHYEDYDGYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEIKPLYEHLH 180

Qy 224 AYVRAKLNNAPSYISPIGCLPAHLGDMWGRFNTNLYSLTVPFGQKPNIDVTDAMVQA 283

Db 181 AYVRPKLMNAPSYISPIGCLPAHLGDMWGRFNTNLYSLTVPFGQKPNIDVTDAMVQA 240

Qy 284 WDAQRIKFAEKFFVSVGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMC 343

Db 241 WDAQRIKFAEKFFVSVGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMC 300

Qy 344 TKVTWDDFLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEMSLSAATPKHLKSI 403

Db 301 TKVTWDDFLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEMSLSAATPKHLKSI 360

Qy 404 GLLSPPDFQEDNETEINFLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMK 463

Db 361 GLLSPPDFQEDNETEINFLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMK 420

Qy 464 REIVGVVEPVPDHEITYCDPASLFHVSNDYSFIRYTRTLVQFQFQALCOAAKHGEGPLHK 523

Db 421 REIVGVVEPVPDHEITYCDPASLFHVSNDYSFIRYTRTLVQFQFQALCOAAKHGEGPLHK 480

Qy 524 CDISNSTEAGQKLFNMLRLKGSBPWTALLENVVGAKNMVRLPLNYFEBLFTWLKDQNKNS 583

Db 481 CDISNSTEAGQKLFNMLRLKGSBPWTALLENVVGAKNMVRLPLNYFEBLFTWLKDQNKNS 540

Qy 584 SFVGNSTDWSPY 595

Db 541 SFVGNSTDWSPY 552

RESULT 9

US-10-158-825-140

; Sequence 140, Application US/10158825

; Patent No. 6900033

; GENERAL INFORMATION:

; APPLICANT: Tom Parry et al.

; TITLE OF INVENTION: Method and Compositions for Modulating ACE-2 Activity

; FILE REFERENCE: PF555

; CURRENT APPLICATION NUMBER: US/10/158,825

; PRIOR FILING DATE: 2002-06-03

; PRIOR APPLICATION NUMBER: 60/294,976

; PRIOR FILING DATE: 2001-06-04

; NUMBER OF SEQ ID NOS: 158

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 140

; LENGTH: 681

; TYPE: PRT

; ORGANISM: homo sapiens

; FEATURE:

; NAME/KEY: MISC FEATURE

; LOCATION: (219)..(219)

; OTHER INFORMATION: Xaa equals any amino acid

; FEATURE:

; NAME/KEY: MISC FEATURE

; LOCATION: (240)..(240)

; OTHER INFORMATION: Xaa equals any amino acid

; FEATURE:

; NAME/KEY: MISC FEATURE

; LOCATION: (499)..(499)

; OTHER INFORMATION: Xaa equals any amino acid

US-10-158-825-140

Query Match 92.4%; Score 2987; DB 2; Length 681;

Best Local Similarity 99.3%; Pred. No. 2.5e-285;

Matches 548; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 44 MNNAGDKWSAFLKEQSTLAQMYPLQEIQLTVKQLQALQOQSSVLSDEKSKRLNTILN 103

Db 1 MNNAGDKWSAFLKEQSTLAQMYPLQEIQLTVKQLQALQOQSSVLSDEKSKRLNTILN 60

Qy 104 TMSITYSTGKVCNPNPQECILLPEGLNEIMANSLDYNERLWAWESWRSVEVGKQLRPLYE 163

Db 61 TMSITYSTGKVCNPNPQECILLPEGLNEIMANSLDYNERLWAWESWRSVEVGKQLRPLYE 120

Qy 164 EYVLKNEMARANHYEDYDGYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEIKPLYEHLH 223

Db 121 EYVLKNEMARANHYEDYDGYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEIKPLYEHLH 180

Qy 224 AYVRAKLNNAPSYISPIGCLPAHLGDMWGRFNTNLYSLTVPFGQKPNIDVTDAMVQA 283

Db 181 AYVRPKLMNAPSYISPIGCLPAHLGDMWGRFNTNLYSLTVPFGQKPNIDVTDAMVQA 240

Qy 284 WDAQRIKFAEKFFVSVGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMC 343

Db 241 WDAQRIKFAEKFFVSVGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMC 300

Qy 344 TKVTWDDFLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEMSLSAATPKHLKSI 403

Db 301 TKVTWDDFLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEMSLSAATPKHLKSI 360

Qy 404 GLLSPPDFQEDNETEINFLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMK 463

Db 361 GLLSPDFQBDNETETINFLKQALITVGTLPFTYMLEKWRWVFKGEIPKQDQNMKKWEMK 420
Qy 464 REIVGVVEVPVPHDETYCDPASLFHVSNDYSFIRYTRTLYQFOQALCOAAKHGSLHK 523
Db 421 REIVGVVEVPVPHDETYCDPASLFHVSNDYSFIRYTRTLYQFOQALCOAAKHGSLHK 480
Qy 524 CDISNSTEAGOKLFNMLRKGKSEPTWTLALENVVGAKNNVRPLLNYFFPLFTWLKDQKN 583
Db 481 CDISNSTEAGOKLFNMLRKGKSEPTWTLALENVVGAKNNVRPLLNYFFPLFTWLKDQKN 540
Qy 584 SPVGMSTWSPY 595
Db 541 SPVGMSTWSPY 552
RESULT 10
US-10-158-847-138
; Sequence 138, Application US/10158847
; Patent No. 6592865
; GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACE-2 Activity
; FILE REFERENCE: PF557
; CURRENT APPLICATION NUMBER: US/10/158,847
; CURRENT FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/295,004
; PRIOR FILING DATE: 2001-06-04
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 138
; LENGTH: 711
; TYPE: PRT
; ORGANISM: homo sapiens
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (219)..(219)
; OTHER INFORMATION: Xaa equals any amino acid
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (240)..(240)
; OTHER INFORMATION: Xaa equals any amino acid
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (499)..(499)
; OTHER INFORMATION: Xaa equals any amino acid
US-10-158-847-138
Query Match 92.4%; Score 2987; DB 2; Length 711;
Best Local Similarity 99.3%; Pred. No. 2.7e-285;
Matches 548; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 44 MNNAGDKWSAFLEKEOSTLAQNYPLQEIQLNTLVKQLQALQONGSSVLSDEKSKRLNTILN 103
Db 1 MNNAGDKWSAFLEKEOSTLAQNYPLQEIQLNTLVKQLQALQONGSSVLSDEKSKRLNTILN 60
Qy 104 TMSTIYSTGKVCNPNPQECILLLEPGLNEIMANSLDYNERLWAWESRSEVKGQLRPLYE 163
Db 61 TMSTIYSTGKVCNPNPQECILLLEPGLNEIMANSLDYNERLWAWESRSEVKGQLRPLYE 120
Qy 164 EYVLKNEANRANHYEDYDWRGDYEVNGVDYDYSRGQLIEDVHTFEETKPLYEHLH 223
Db 121 EYVLKNEANRANHYEDYDWRGDYEVNGVDYDYSRGQLIEDVHTFEETKPLYEHLH 180
Qy 224 AYVRKLNANPYSYISPIGCLPAHLGDMWGRFTNLSLTPFGQKPNIDVTDAMVQA 283
Db 181 AYVRKLNANPYSYISPIGCLPAHLGDMWGRFTNLSLTPFGQKPNIDVTDAMVQA 240
Qy 284 WDAQRIFKEAEKFFVSUGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMC 343
Db 241 WDAQRIFKEAEKFFVSUGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMC 300
Qy 344 TKVTMDDFLTAHMGHIIQYDMAVAAQPFLLRNGANEGFHEAVGBIMSLSAATPKHLKSI 403

Db 301 TKVTMDDFLTAHMGHIIQYDMAVAAQPFLLRNGANEGFHEAVGBIMSLSAATPKHLKSI 360
Qy 404 GLLSPDFQBDNETETINFLKQALITVGTLPFTYMLEKWRWVFKGEIPKQDQNMKKWEMK 463
Db 361 GLLSPDFQBDNETETINFLKQALITVGTLPFTYMLEKWRWVFKGEIPKQDQNMKKWEMK 420
Qy 464 REIVGVVEVPVPHDETYCDPASLFHVSNDYSFIRYTRTLYQFOQALCOAAKHGSLHK 523
Db 421 REIVGVVEVPVPHDETYCDPASLFHVSNDYSFIRYTRTLYQFOQALCOAAKHGSLHK 480
Qy 524 CDISNSTEAGOKLFNMLRKGKSEPTWTLALENVVGAKNNVRPLLNYFFPLFTWLKDQKN 583
Db 481 CDISNSTEAGOKLFNMLRKGKSEPTWTLALENVVGAKNNVRPLLNYFFPLFTWLKDQKN 540
Qy 584 SPVGMSTWSPY 595
Db 541 SPVGMSTWSPY 552
RESULT 11
US-10-158-825-138
; Sequence 138, Application US/10158825
; Patent No. 6900033
; GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACE-2 Activity
; FILE REFERENCE: PF555
; CURRENT APPLICATION NUMBER: US/10/158,825
; CURRENT FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/294,976
; PRIOR FILING DATE: 2001-06-04
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 138
; LENGTH: 711
; TYPE: PRT
; ORGANISM: homo sapiens
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (219)..(219)
; OTHER INFORMATION: Xaa equals any amino acid
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (240)..(240)
; OTHER INFORMATION: Xaa equals any amino acid
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (499)..(499)
; OTHER INFORMATION: Xaa equals any amino acid
US-10-158-825-138
Query Match 92.4%; Score 2987; DB 2; Length 711;
Best Local Similarity 99.3%; Pred. No. 2.7e-285;
Matches 548; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 44 MNNAGDKWSAFLEKEOSTLAQNYPLQEIQLNTLVKQLQALQONGSSVLSDEKSKRLNTILN 103
Db 1 MNNAGDKWSAFLEKEOSTLAQNYPLQEIQLNTLVKQLQALQONGSSVLSDEKSKRLNTILN 60
Qy 104 TMSTIYSTGKVCNPNPQECILLLEPGLNEIMANSLDYNERLWAWESRSEVKGQLRPLYE 163
Db 61 TMSTIYSTGKVCNPNPQECILLLEPGLNEIMANSLDYNERLWAWESRSEVKGQLRPLYE 120
Qy 164 EYVLKNEANRANHYEDYDWRGDYEVNGVDYDYSRGQLIEDVHTFEETKPLYEHLH 223
Db 121 EYVLKNEANRANHYEDYDWRGDYEVNGVDYDYSRGQLIEDVHTFEETKPLYEHLH 180
Qy 224 AYVRKLNANPYSYISPIGCLPAHLGDMWGRFTNLSLTPFGQKPNIDVTDAMVQA 283
Db 181 AYVRKLNANPYSYISPIGCLPAHLGDMWGRFTNLSLTPFGQKPNIDVTDAMVQA 240
Qy 284 WDAQRIFKEAEKFFVSUGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMC 343

Db 241 WDAQRIFKAEKFFVSVGLPNNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLM 300
QY 344 TKVTMDPDLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGIMSLSAATPKHLKSI 403
Db 301 TKVTMDPDLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGIMSLSAATPKHLKSI 360
QY 404 GLISPDQDNTEINFLKQALTIIVGTLPFTYMLEKRWVFKGEIPKQOMKQWENK 463
Db 361 GLISPDQDNTEINFLKQALTIIVGTLPFTYMLEKRWVFKGEIPKQOMKQWENK 420
QY 464 REIVGVVPEVPHDETCDPASLFHVSNDYSFIRYTRTLYQFQFALCOAAKHGGLHK 523
Db 421 REIVGVVPEVPHDETCDPASLFHVSNDYSFIRYTRTLYQFQFALCOAAKHGGLHK 480
QY 524 CDISNSTEAGQKLFNMLRGKSEPTALLENVVGAKNMNVRPLNYPFLTWLKDQNK 583
Db 481 CDISNSTEAGQKLFNMLRGKSEPTALLENVVGAKNMNVRPLNYPFLTWLKDQNK 540
QY 584 SFVGMSTWSPY 595
Db 541 SFVGMSTWSPY 552

RESULT 12
US-08-481-626-2
; Sequence 2, Application US/08481626
; Patent No. 5801040
; GENERAL INFORMATION:
; APPLICANT: Soubrier, Florent
; APPLICANT: Ahenc-Gelas, Francois
; APPLICANT: Hubert, Christine
; APPLICANT: Corvol, Pierre
; TITLE OF INVENTION: Nucleic Acid Coding for the Human
; TITLE OF INVENTION: Testicular Angiotensin Converting Enzyme (ACE) and its
; TITLE OF INVENTION: Uses, Especially for the In Vitro Screening for this
; TITLE OF INVENTION: Enzyme in the Organism
; NUMBER OF SEQUENCES: 3
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Finnegan, Henderson, Farabow, Garrett &
; ADDRESS: 1300 I Street, N.W., Suite 700
; CITY: Washington
; STATE: D.C.
; ZIP: 20005-3315
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/481,626
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/656,183
; FILING DATE: 04-MAR-1991
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: FR 89-09062
; FILING DATE: 05-JUL-1989
; ATTORNEY/AGENT INFORMATION:
; NAME: Meyers, Kenneth J.
; REGISTRATION NUMBER: 25,146
; REFERENCE/DOCKET NUMBER: 04958-0006-02000
; TELEPHONE: 202-408-4000
; TELEFAX: 202-408-4400
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 732 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear

; MOLECULE TYPE: protein
US-08-481-626-2
Query Match 41.3%; Score 1335; DB 1; Length 732;
Best Local Similarity 41.9%; Pred. No. 2,4e-122;
Matches 254; Conservative 116; Mismatches 202; Indels 34; Gaps 9;
QY 2 TIEQAKTFLDKFHEAEDLPYQSSLASNNYNTNITEB-----NVQNNMNAQDKWSA 53
Db 70 TDEAASKFVEYDRTSQVWVNEVAEANNYNTNITTTSTKILLQKNQIAHNT----- 123
QY 54 FLKEOSTLAQMYPLQEIQLNLTQKLOALQNGSSVLTSEDKSKRLNTILNTMTSTYSTCK 113
Db 124 --LKYGTQARKFDVNLQNTTIKRIIKVQDLERAAALPAQELSEYNKILLDMETTYSVAT 181
QY 114 VCNPNPOCEILLPEGLNEIMANSIDYNERLWAMESEVKGQKRLPYEYVVLKNEWA 173
Db 182 VCHPNG--SCLEQEPDLTNWATSEKYEDLLWAMEGWDKAGRAILQYFYPKVELINQAA 239
QY 174 RANHYEDYGDYWRGDIYVNGVDYDSRGQILIEDVEHTFEEIKPLYEHLHAYVRAKLMA 233
Db 240 RLNGYVDAGDSWRSWYETPSLE-----QDLERLFQELQPLLYLNLHAYVREALHRH 289
QY 234 Y-PSYISPIGCLPAHLGDMGRFWNLXSLTVPFGQKFNIDVTDAMVDQAWDAQRIKKE 292
Db 290 YGAQHINLEGPIPAHLGDMWAQTNSTYDLYVPPSPASMDTTEAMLKQGWTPRRMFKE 349
QY 293 AEKFFSVGLPNNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKG-DPRILMCTKVTMDDF 351
Db 350 ADDFTSLGLLPVPEPFWNKSMLEKPTDGRVVCHASANDFYNGKDFRIKQCTTVNLEDL 409
QY 352 LTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGIMSLSAATPKHLKSLGLSPDFQ 411
Db 410 VVAHHEMGHIQYFMQYKDLPLVALREGANPGFHEAGIDVLALSVPKHLHSLNLLSSEGG 469
QY 412 EDNETEINFLKQALTIIVGTLPFTYMLEKRWVFKGEIPKQOMKQWENKREIVGVVE 471
Db 470 SD-EHDINFLMKVALDKIAFPFSYLVQWRVDFGSIITKENYQEWWSLRLKYQGLCP 528
QY 472 VVPHDETCDPASLFHVSNDYSFIRYTRTLYQFQFALCOAAKHGGLHKCDISNSTE 531
Db 529 VVPRTOGDFDCAKEHIPSSVPIRYFYFIIQFQFHEALCOAGHTGFLHKCDIYQSK 588
QY 532 AQOKLFNMLRGKSEPTALLENVVGAKNMNVRPLNYPFLTWLKDQNK--NSFVGH- 588
Db 589 AGQRLATAMKLGFSRPPWPEAMQLITQPNMSASAMLSYFKPLDLLMLRTENELHGEKLGWP 648
QY 589 STDWSP 594
Db 649 QYNWTP 654

RESULT 13
US-08-989-299-4
; Sequence 4, Application US/08989299
; Patent No. 6194556
; GENERAL INFORMATION:
; APPLICANT: Acton, Susan L.
; APPLICANT: Robinson, Keith E.
; TITLE OF INVENTION: ANGIOTENSIN CONVERTING ENZYME HOMOLOG
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: FOLEY, HOAG & ELIOT LLP
; CITY: Boston
; STATE: MA
; COUNTRY: USA
; ZIP: 02109-2170
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS


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; Patent No. 6884771
; GENERAL INFORMATION:
; APPLICANT: Acton, Susan L. et al.
; TITLE OF INVENTION: ANGIOTENSIN CONVERTING ENZYME HOMOLOG AND THERAPEUTIC
; TITLE OF INVENTION: DIAGNOSTIC USES THEREFOR
; FILE REFERENCE: MNI-132CP3
; CURRENT APPLICATION NUMBER: US/09/635,501
; CURRENT FILING DATE: 2000-08-09
; PRIOR APPLICATION NUMBER: 09/407,427
; PRIOR FILING DATE: 1999-09-29
; PRIOR APPLICATION NUMBER: 09/163,648
; PRIOR FILING DATE: 1998-09-30
; PRIOR APPLICATION NUMBER: 08/989,299
; PRIOR FILING DATE: 1997-12-11
; NUMBER OF SEQ ID NOS: 107
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 4
; LENGTH: 732
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-635-501-4

Query Match 41.3%; Score 1335; DB 2; Length 732;
Best Local Similarity 41.9%; Pred. No 2.4e-122;
Matches 254; Conservative 116; Mismatches 202; Indels 34; Gaps 9;

QY 2 TIEQAKTFLDKFNBEADLFYQSSLASWNYNTNITEE-----NVQNMNAGDKWSA 53
DB 70 TDEAEASKFVEEDRTSQVWNEVEANWNYNTNITETSKILLQKNMQIANHT----- 123
QY 54 FLKEQSTLAQMYPLQIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGK 113
DB 124 --LKYGTQARKPDVNLQNTIKRIIKYQDLERAAAPAELEENKILLDMETTYSVAT 181
QY 114 VCNPDNPQECLELLEGLNEIMANSLDYNERLWAEWSRSVKGQRLAPLYEEYVVLKNEMA 173
DB 182 VCHPNG--SCLQLEPDLTNMATSRKYEDLLWAEGRDAGRAILOFTPKYVELINQAA 239
QY 174 RANHYEDYGDYWRGDEVNGVDGYDYSRGOLIEDVHTFEIKPLYEHLHAYVRAKLMA 233
DB 240 RLNGYVDAGDSRSMYETPSLE-----QDLERLFQELQPLYLNLHAYVRRALHRH 289
QY 234 Y-PSYISPTGCLPAHLIGDMGRFNTNLSLTVPPGQKPNIDVTDAMVDQAWDAQRIFK 292
DB 290 YGAQHINLEGPIPAHLIGNMWAOQWTSNIYDLVVPSPAPSMDTTEAMLKQGWTPRRMPKE 349
QY 293 AEKFFVSVGLPNNMQGFWNSMLTDPGNVQKAVCHPTANDLGKG-DPRILMCTKVTMDDF 351
DB 350 ADDFTSLGLLPVPPEFWNKSMLDKPTDGRVGVCHASAWDFYNGKDFRIKQCTTVNLEDL 409
QY 352 LTAHEMIGHIQYDMAYAAQPFLLRNGANEGFEAVGEIMSLSAATPKHLKSLGLLSPDFQ 411
DB 410 VVAHEMIGHIQYPMQYKDLPLVALREGANPGFHEAIGDVLALSVSPTPKHLHSLNLSSEGG 469
QY 412 EDNETEINFLLKQALITVGLTPPTMYLEKRWNVFKGEIPKQOMMKKWMKREIVGVVE 471
DB 470 SD-EHDINFLLKMAKDIAFIPESYLVLDQWRWRVFDGSIKENYNGEWSRLKYLKQGLCP 528
QY 472 PVPHDITYCDPASLHVNDYSIRYRTLTQFQFQALCOAAKHEGPLHKCDISNSTE 531
DB 529 PVPTQGFDFGAKFHIPSSVPIRYVFSFIQFQFHEALCOAAGTGPLHKCDIYQSK 588
QY 532 AGQKLFNMLRLKGSSEPTWLTALENVGAKNNVRPLINYPEPLFTWLKQNK--NSFVGW- 588
DB 589 AGQRLATAMKLGFSRPWPEAMQLITGQPNNSASAMLSYFKPLDLDLRTENELHGEKLGWP 648
QY 589 STDWSP 594
DB 649 QYNWTP 654
```

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GenCore version 5.1.7
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OM protein - protein search, using sw model

Run on: March 28, 2006, 11:02:42 ; Search time 135.885 Seconds
(without alignments)
1923.914 Million cell updates/sec

Title: US-10-659-000-4
Perfect score: 3231
Sequence: 1 STIEQAKTFLDKFNHEAD.....WLKQNKNSFVGNSTDMSPY 595

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 2443163 seqs, 439378781 residues

Total number of hits satisfying chosen parameters: 2443163

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_21.*
1: Geneseqp1980s:*
2: Geneseqp1990s:*
3: Geneseqp2000s:*
4: Geneseqp2001s:*
5: Geneseqp2002s:*
6: Geneseqp2003as:*
7: Geneseqp2003bs:*
8: Geneseqp2004s:*
9: Geneseqp2005s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3231	100.0	595	8 ADO15670	Ado15670 Human ACE
2	3231	100.0	771	8 ABM85139	Abm85139 Human dia
3	3231	100.0	805	3 AAY84562	Aay84562 A human a
4	3231	100.0	805	3 AAY67310	Aay67310 Human MPR
5	3231	100.0	805	4 AAB48095	Aab48095 Human Zac
6	3231	100.0	805	4 AAY72667	Aay72667 Human ang
7	3231	100.0	805	5 AAE20353	Aae20353 Human ACE
8	3231	100.0	805	5 ABG77023	Abg77023 Human ang
9	3231	100.0	805	5 ABG77023	Abg77023 Human ang
10	3231	100.0	805	5 AAU99701	Aau99701 Human ang
11	3231	100.0	805	6 ABU07731	Abu07731 Human zin
12	3231	100.0	805	6 ADA03344	Ada03344 Human ang
13	3231	100.0	805	6 ABR56712	Abr56712 Human ACE
14	3231	100.0	805	7 ADL95395	Adl95395 Human ang
15	3231	100.0	805	7 ADL95494	Adl95494 Human ang
16	3231	100.0	805	8 ADH51357	Adh51357 Human ang
17	3231	100.0	805	8 ADO55144	Ado55144 Protein #
18	3231	100.0	805	9 ADZ27233	Adz27233 Human ACE
19	3227	99.6	805	7 ADC38728	Adc38728 Human sec
20	3227	99.6	702	9 ADZ27234	Adz27234 Soluble h
21	2998.5	92.8	768	8 ABM85140	Abm85140 Human dia
22	2987	92.4	681	6 ADA03342	Ada03342 Human ang
23	2987	92.4	681	6 ABR56711	Abr56711 Human ACE
24	2987	92.4	711	4 AAU09092	Aau09092 Novel hum

25	2987	92.4	711	6 ADO03340	Ada03340 Human ang
26	2987	92.4	711	6 ABR56709	Abr56709 Human ACE
27	2987	89.7	555	4 AAU12207	Aau12207 Human PRO
28	2987	89.7	555	6 ABO17651	Abu17651 Novel hum
29	2987	89.7	555	6 ABU80905	Abu80905 Human PRO
30	2987	89.7	555	6 ABU66605	Abu66605 Human PRO
31	2987	89.7	555	6 ABU59686	Abu59686 Novel sec
32	2987	89.7	555	6 ABO24876	Abu24876 Human sec
33	2987	89.7	555	6 ABU66881	Abu66881 Human sec
34	2987	89.7	555	6 ADA45591	Ada45591 Novel hum
35	2987	89.7	555	6 ADA76022	Ada76022 Human PRO
36	2987	89.7	555	6 ADA18672	Ada18672 Human PRO
37	2987	89.7	555	6 ADA61295	Ada61295 Homo sapi
38	2987	89.7	555	6 ADB19080	Adb19080 Novel hum
39	2987	89.7	555	6 ADB27621	Adb27621 Human PRO
40	2987	89.7	555	6 ADA86100	Ada86100 Novel hum
41	2987	89.7	555	6 ADB15664	Adb15664 Human PRO
42	2987	89.7	555	6 ADA47450	Ada47450 Human PRO
43	2987	89.7	555	6 ADA67245	Ada67245 Human PRO
44	2987	89.7	555	6 ADB30252	Adb30252 Human PRO
45	2987	89.7	555	6 ADA85548	Ada85548 Novel hum

ALIGNMENTS

RESULT 1
ADO15670
ID ADO15670 standard; protein; 595 AA.
XX
AC ADO15670;
XX
DT 01-JUL-2004 (first entry)
XX
DE Human ACE2, SEQ ID 4.
XX
KW protein co-ordinate data; crystal;
KW angiotensin-converting enzyme-related carboxypeptidase; ACE2;
KW angiotensin-converting enzyme; ACE; enzyme.
XX
OS Homo sapiens.
XX
PN WO2004023270-A2.
XX
PD 18-MAR-2004.
XX
PF 09-SEP-2003; 2003WO-US028374.
XX
PR 09-SEP-2002; 2002US-0410010P.
XX
(MILL-) MILLENIUM PHARM INC.
XX
PI Pantollano MW, Ryan MD, Staker BL, Prasad GS, Tang J, Menon SP;
PI Towler PS, Williams DH, Fisher M;
XX
DR WPI; 2004-315606/29.
XX
PT Crystal of angiotensin-converting enzyme-related carboxypeptidase or its
PT homolog, useful for detecting compounds e.g. ligands capable of binding
PT to angiotensin-converting enzyme-related carboxypeptidase.
XX
PS Claim 6; Fig 4; 539pp; English.
XX
CC The present invention relates to a crystal (I) comprising an angiotensin-
CC converting enzyme-related carboxypeptidase (ACE2) or its homolog. (I) is
CC useful for detecting chemical compounds such as ligand, antagonist,
CC agonist, inhibitor, antibody, peptide, protein or drug having capability
CC of binding to the active site of the ACE2 protein. The present sequence
CC is human ACE2, used to illustrate the invention.
XX
SQ Sequence 595 AA;

Query Match 100.0%; Score 3231; DB 8; Length 595;

Best Local Similarity 100.0%; Pred. No. 3.8e-288;		Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
QY	1	STIEEQAKTFDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWAFLEKEQST	60
Db	1	STIEEQAKTFDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWAFLEKEQST	60
QY	61	LAQMYPLQEIQNLTVKQLQALQQNGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP	120
Db	61	LAQMYPLQEIQNLTVKQLQALQQNGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP	120
QY	121	QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEEYVVLKNEMARANHYED	180
Db	121	QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEEYVVLKNEMARANHYED	180
QY	181	YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRACLNNAYPSYISP	240
Db	181	YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRACLNNAYPSYISP	240
QY	241	IGCLPAHLIGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSU	300
Db	241	IGCLPAHLIGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSU	300
QY	301	GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVTMDDFLTAAHEMGH	360
Db	301	GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVTMDDFLTAAHEMGH	360
QY	361	IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGILSLSPDFQEDNTEINF	420
Db	361	IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGILSLSPDFQEDNTEINF	420
QY	421	LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQOMKWKWEMKREIVGVVEVPVPHDETYC	480
Db	421	LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQOMKWKWEMKREIVGVVEVPVPHDETYC	480
QY	481	DPASLFHVSNDYSFIRYTRTLTYQFQFQALCQAAKHGEGPLHKCDISNSTEAGQKLFNNL	540
Db	481	DPASLFHVSNDYSFIRYTRTLTYQFQFQALCQAAKHGEGPLHKCDISNSTEAGQKLFNNL	540
QY	541	RLGKSEPTIALENVVGAKNMVRPLLNYFEPLFTWLKDQNKNSFVGNSTDNSPY	595
Db	541	RLGKSEPTIALENVVGAKNMVRPLLNYFEPLFTWLKDQNKNSFVGNSTDNSPY	595

RESULT 2

ID	ABM85139	standard; protein; 771 AA.
XX	AC	ABM85139;
XX	AC	ABM85139;
DT	18-NOV-2004	(first entry)
DE	Human	diagnostic and therapeutic pprotein SEQ ID NO:5388.
XX	gene therapy; human	diagnostic and therapeutic polynucleotide; dithp.
XX	Homo sapiens.	
XX	WO2004023973-A2.	
XX	25-MAR-2004.	
PF	12-SEP-2003;	2003WO-US028227.
XX	12-SEP-2002;	2002US-0410259P.
PR	12-SEP-2002;	2002US-0410260P.
XX	(INCY-)	INCYTE CORP.
XX	Schmidt JP,	Wright RJ, Bruns CM, Marjanovic MM, Shen F;
PI	Harthshorne TA,	Suchorolski MT, Altus CM, Pitts SJ, Elder LV;
PI	Mooney BM,	Delegane AM, Panesar IS, Banville SC, Reddy TP;
PI	Stevens KA,	Blanchard JL, Panzer SR, Wang X, Au AP, Gerstin EH;

PI	Peralta CH,	Anderson SB, Rioux P, Shen EJ, Wu MC, Stuve LL;
PI	Lagace RE,	Spiro PA, Stewart EA, Wingrove J, Vitt UA, Kirtson ES;
PI	Xu Y,	Kwong M, Policky JL, Hurwitz BL, Ma Y, Jackson JL, Gietzen D;
PI	Patury S,	Shi X, Suarez CJ;
DR	WPI:	2004-329368/30.
DR	N-PSDB;	ACN43791.
XX		
PT	New	diagnostic and therapeutic polynucleotides and polypeptides, useful
PT	in	diagnosing a condition, disease or disorder associated with human
PT	molecules,	e.g. autoimmune or inflammatory disorders, in gene therapy or
PT	in	gene mapping.
XX		
PS	Claim 27;	Page; 190pp; English.
XX		
CC	The	invention relates to novel diagnostic and therapeutic polynucleotides
CC	selected	from one of the 2722 sequences defined in the specification. A
CC	polynucleotide	of the invention may have a use in gene therapy. The human
CC	diagnostic	and therapeutic polynucleotides (dithp) or polypeptides may be
CC	used	to diagnose a particular condition, disease or disorder associated
CC	with	human molecules, e.g. cell proliferative disorders,
CC	autoimmune/inflammatory	disorder, developmental disorder, endocrine
CC	disorder,	neurological disorders, gastrointestinal disorders, or
CC	infections	caused by virus, bacteria, fungi or parasite. The dithp
CC	molecules	may also be used in genetic mapping, in identifying individuals
CC	from	minute biological samples, in detecting single nucleotide
CC	polymorphisms,	as molecular weight markers, and for somatic or germline
CC	gene	therapy. The present sequence represents a dithp protein of the
CC	invention.	Note: the sequence data for this patent is not represented in
CC	the	printed specification, but was obtained in electronic format directly
CC	from	WIPO at www.wipo.int/pct/en/sequences/listing.htm
XX		
SQ	Sequence 771 AA;	
	Query Match	100.0%; Score 3231; DB 8; Length 771;
	Best Local Similarity	100.0%; Pred. No. 5.7e-288;
	Matches 595;	Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1	STIEEQAKTFDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWAFLEKEQST 60
Db	19	STIEEQAKTFDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWAFLEKEQST 78
QY	61	LAQMYPLQEIQNLTVKQLQALQQNGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 120
Db	79	LAQMYPLQEIQNLTVKQLQALQQNGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 138
QY	121	QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEEYVVLKNEMARANHYED 180
Db	139	QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEEYVVLKNEMARANHYED 198
QY	181	YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRACLNNAYPSYISP 240
Db	199	YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRACLNNAYPSYISP 258
QY	241	IGCLPAHLIGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSU 300
Db	259	IGCLPAHLIGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSU 318
QY	301	GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVTMDDFLTAAHEMGH 360
Db	319	GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVTMDDFLTAAHEMGH 378
QY	361	IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGILSLSPDFQEDNTEINF 420
Db	379	IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSGILSLSPDFQEDNTEINF 438
QY	421	LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQOMKWKWEMKREIVGVVEVPVPHDETYC 480
Db	439	LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQOMKWKWEMKREIVGVVEVPVPHDETYC 498
QY	481	DPASLFHVSNDYSFIRYTRTLTYQFQFQALCQAAKHGEGPLHKCDISNSTEAGQKLFNNL 540
Db	499	DPASLFHVSNDYSFIRYTRTLTYQFQFQALCQAAKHGEGPLHKCDISNSTEAGQKLFNNL 558

OY 541 RLKSEPTLALENVGAKNMVRLPLNTYFEPFLTWLKDQNKNSFVGWSTDWSPY 595
 DB 559 RLKSEPTLALENVGAKNMVRLPLNTYFEPFLTWLKDQNKNSFVGWSTDWSPY 613

RESULT 3

AA84562
 ID AAY84562 standard; protein; 805 AA.

XX AC AAY84562;

DT 25-JUL-2000 (first entry)

XX A human angiotensin converting enzyme-2 (ACE-2) protein.

KW Human; angiotensin converting enzyme-2; ACE-2; angiotensin I; Ang. (1-9);
 KW blood pressure; hypertension; congestive heart failure; atherosclerosis;
 KW chronic heart failure; acute heart failure; myocardial infarction;
 KW renal failure.

XX Homo sapiens.

XX Location/Qualifiers

FT Key

FT Peptide

FT 1..18

FT /note= "signal sequence"

FT Domain

FT 19..740

FT /note= "extracellular domain"

FT Domain

FT 374..378

FT /note= "minimal zinc binding domain"

FT Domain

FT 741..765

FT /note= "transmembrane domain"

FT Domain

FT 766..805

FT /note= "cytoplasmic domain"

XX WO200018899-A2.

XX 06-APR-2000.

XX 29-SEP-1999; 99WO-US022976.

XX 30-SEP-1998; 98US-00163648.

XX (MILL-) MILLENNIUM PHARM INC.

XX Acton LS, Robison KE, Hsieh FY;

XX WPI; 2000-293140/25.

XX N-PSDB; AAA12764.

XX Isolated nucleic acid encoding angiotensin converting enzyme-2 (ACE-2)

XX polypeptide useful for detecting an ACE-2 therapeutic for treating

XX hypertension, congestive heart failure, myocardial infarction,

XX atherosclerosis and renal failure.

XX Claim 2; Fig 1; 138pp; English.

XX The present sequence represents a human angiotensin converting enzyme-2
 CC (ACE-2). ACE-2 is expressed predominantly in kidneys and testis. The
 CC sequence of the full length ACE-2 cDNA was determined from a clone
 CC obtained from a cDNA library prepared from mRNA of a human heart of a
 CC subject who had congestive heart failure. ACE-2 has significant sequence
 CC homologies with ACE enzymes, and has also been shown to hydrolyse
 CC angiotensin I into Ang. (1-9). The ACE-2 therapeutics are used to treat
 CC blood pressure related diseases and conditions, such as hypertension,
 CC congestive heart failure, chronic heart failure, acute heart failure,
 CC myocardial infarction, atherosclerosis and renal failure

XX Sequence 805 AA;

XX Query Match

XX Best Local Similarity 100.0%; Score 3231; DB 3; Length 805;

XX Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 STIERQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKSAFLKEOST 60
 DB 19 STIERQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKSAFLKEOST 78
 OY 61 LAQMPLOBIQNLTVKLOLQALQQNGSSVLSKSKRLNTILNTMTSTYTGKVCNPNP 120
 DB 79 LAQMPLOBIQNLTVKLOLQALQQNGSSVLSKSKRLNTILNTMTSTYTGKVCNPNP 138
 OY 121 QECLELLEPGLNEIMANSLDYNERLWAWESRSEVGKQLRPLYEYVVLKNEWARANHYED 180
 DB 139 QECLELLEPGLNEIMANSLDYNERLWAWESRSEVGKQLRPLYEYVVLKNEWARANHYED 198
 OY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEIKPLYEHLHAYVRAKLMNAYPSYIS 240
 DB 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVEHTPEEIKPLYEHLHAYVRAKLMNAYPSYIS 258
 OY 241 IGCLPAHLIGDMWGRFTNLVSLVFPQKPNIDVTDAMVDQAWDAQRIKFAEKFFVS 300
 DB 259 IGCLPAHLIGDMWGRFTNLVSLVFPQKPNIDVTDAMVDQAWDAQRIKFAEKFFVS 318
 OY 301 GLPNNTQGFWENSMLTDFGNVQKAVCHPTAMDGLKGFRIILMCTKTVMDDFLTAHHEM 360
 DB 319 GLPNNTQGFWENSMLTDFGNVQKAVCHPTAMDGLKGFRIILMCTKTVMDDFLTAHHEM 378
 OY 361 IOYDMAYAAQPPLLRNGANEGFHEAVGIMSLSAATPKHLKSIIGLLSPDFQDNTEINF 420
 DB 379 IOYDMAYAAQPPLLRNGANEGFHEAVGIMSLSAATPKHLKSIIGLLSPDFQDNTEINF 438
 OY 421 LLKQALTIIVGTLPTFTMLEKRWVMVFKGEIPKQDMKKWEMKREIVGVVVPVPHDETYC 480
 DB 439 LLKQALTIIVGTLPTFTMLEKRWVMVFKGEIPKQDMKKWEMKREIVGVVVPVPHDETYC 498
 OY 481 DPASLFHVSNDYSFIRYTRTYLQFQFQALCQAAKHEGLKCDISNSTEAGQKLFNNL 540
 DB 499 DPASLFHVSNDYSFIRYTRTYLQFQFQALCQAAKHEGLKCDISNSTEAGQKLFNNL 558
 OY 541 RLKSEPTLALENVGAKNMVRLPLNTYFEPFLTWLKDQNKNSFVGWSTDWSPY 595
 DB 559 RLKSEPTLALENVGAKNMVRLPLNTYFEPFLTWLKDQNKNSFVGWSTDWSPY 613

RESULT 4

AA67310

ID AAY67310 standard; protein; 805 AA.

XX AC AAY67310;

XX 11-APR-2000 (first entry)

XX Human MPROT15 amino acid sequence #1.

XX MPROT15; treatment; hypertension; human; myocardial disease; apoplexy;
 KW heart disease; apoplexy; heart disease; nervous denaturation; hormone;
 KW Alzheimer's disease; cytokine.

XX Homo sapiens.

XX JPL11318472-A.

XX 24-NOV-1999.

XX 22-JAN-1999; 99JP-00014949.

XX 13-MAY-1998; 98GB-00010373.

XX 18-AUG-1998; 98GB-00018009.

XX (SMIK) SMITHKLINE BEECHAM PLC.

XX WPI; 2000-109268/10.

XX N-PSDB; AA259465.

XX MPROT15 polypeptide and MPROT15 polynucleotides - useful for the

PT treatment of hypertension, myocardial diseases, apoplexy, heart diseases,
XX nervous denaturation, Alzheimer's disease etc.

PS Claim 1; Page 15; 22pp; Japanese.

XX This is amino acid sequence #1 of human MPROT15. The MPROT15
CC polynucleotide and polypeptide sequences can be used for the treatment of
CC hypertension, myocardial diseases, apoplexy, heart diseases, nervous
CC denaturation, Alzheimer's disease and diseases related to the processing
CC of peptide hormones and cytokines

XX SQ Sequence 805 AA;

Query Match 100.0%; Score 3231; DB 3; Length 805;
Best Local Similarity 100.0%; Pred. No. 6.1e-288;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLEKEQST 60
Db |||||
19 STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLEKEQST 78
QY 61 LAQMYPLOBIQNLTVKQLQALQONGSSVLSDEKSKRLNTILNTWSTIYSTGKVCNPNP 120
Db |||||
79 LAQMYPLOBIQNLTVKQLQALQONGSSVLSDEKSKRLNTILNTWSTIYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEMARANHYED 180
Db |||||
139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEMARANHYED 198
QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEETKPLYEHLHAYVRAKLMNAYSISYSP 240
Db |||||
199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEETKPLYEHLHAYVRAKLMNAYSISYSP 258
QY 241 IGCLPAHLGDMWGRFWNLVSLTVFPGQKPNIDVTDAMVDQAWDAQRIKFAEKEFFVSV 300
Db |||||
259 IGCLPAHLGDMWGRFWNLVSLTVFPGQKPNIDVTDAMVDQAWDAQRIKFAEKEFFVSV 318
QY 301 GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMGGH 360
Db |||||
319 GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMGGH 378
QY 361 IQYDMAYAAQPFLLRNGANGFHEAEGEIMSLAATPKHLKIGLLSPDFQEDNETEINF 420
Db |||||
379 IQYDMAYAAQPFLLRNGANGFHEAEGEIMSLAATPKHLKIGLLSPDFQEDNETEINF 438
QY 421 LKQALTIIVGTLPFTYMLEKRWMMVFKGIPKQDNKKWEMKREIVGVVPEVPHDETTC 480
Db |||||
439 LKQALTIIVGTLPFTYMLEKRWMMVFKGIPKQDNKKWEMKREIVGVVPEVPHDETTC 498
QY 481 DPASLPHVNDYSFIRYVTRTYQFQBALQAAKHEGPHKCDISNSTEAGOKLFNNML 540
Db |||||
499 DPASLPHVNDYSFIRYVTRTYQFQBALQAAKHEGPHKCDISNSTEAGOKLFNNML 558
QY 541 RLKGSPEWTLALENVVGAKNMVRPLLNYFEPFLTWLKDQNKNSFVGNSTWSPY 595
Db |||||
559 RLKGSPEWTLALENVVGAKNMVRPLLNYFEPFLTWLKDQNKNSFVGNSTWSPY 613

RESULT 5
AAB48095

ID AAB48095 standard; protein; 805 AA.

XX AAB48095;

XX 19-MAR-2001 (first entry)

DE Human Zace2 protein.

XX Zace2; metalloenzyme; angiotensin-converting enzyme; ACE; fertility;
KW zinc metalloproteinase; blood pressure; zinc protease; hypertension;
KW ventricular systolic dysfunction; renal impairment; heart failure;
KW scleroderma renal crisis; atherosclerosis; antiinflammatory; human;
KW antiarthritic; bradykinin inactivator.

XX Homo sapiens.
XX WO200070032-A1.
XX 23-NOV-2000.
XX 03-MAY-2000; 2000WO-US011932.
XX 13-MAY-1999; 99US-00311482.
XX 27-AUG-1999; 99US-00384706.
XX (ZYMO) ZYMOGENETICS INC.
XX Piddington CS, Petrie CR, Shoemaker KE, Bishop PD;
XX WPI; 2001-025018/03.
XX N-PSDB; AAC84366, AAC84367.
XX Angiotensin-converting enzyme, Zace2, useful for treating inflammatory
XX bowel disease, e.g. Crohn's disease and ulcerative colitis, or diseases
XX associated with inflammation such as arthritis and enterocolitis.
XX Example 1; Page 95-100; 125pp; English.

XX The invention relates to the metalloenzyme Zace2. Zace2, an angiotensin-
XX converting enzyme is a zinc metalloproteinase that plays roles in blood
XX pressure regulation and fertility. Zace2 can be expressed by standard
XX recombinant methodology. Zace2 polypeptides are useful for treating an
XX inflammatory bowel disease (e.g. Crohn's disease and ulcerative colitis),
XX diseases associated with inflammation like arthritis and enterocolitis,
XX as targets for identifying modulators of zinc protease activity, for
XX screening or identifying new angiotensin-converting enzyme (ACE)
XX inhibitors, and as a basis for rational drug design for inhibitory
XX molecules. The nucleic acids can be used to detect the expression of a
XX Zace2 gene in a biological sample, as probes for in vivo diagnosis and
XX for detecting and localizing Zace2 gene expression in tissue samples, to
XX determine whether a subject's chromosomes contain a mutation in the Zace2
XX gene, and to detect aberrations associated with the Zace2 locus.
XX Inhibitors of ACE are used for treating hypertension of various
XX conditions, including left ventricular systolic dysfunction, progressive
XX renal impairment, scleroderma renal crisis, congestive heart failure due
XX to dysfunction, and treatment of atherosclerosis. Zace2 agonists may be
XX used to treat infertility while Zace2 antagonists are used for inducing
XX infertility. The present sequence represents the human Zace2 protein

XX SQ Sequence 805 AA;

Query Match 100.0%; Score 3231; DB 4; Length 805;
Best Local Similarity 100.0%; Pred. No. 6.1e-288;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLEKEQST 60
Db |||||
19 STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLEKEQST 78
QY 61 LAQMYPLOBIQNLTVKQLQALQONGSSVLSDEKSKRLNTILNTWSTIYSTGKVCNPNP 120
Db |||||
79 LAQMYPLOBIQNLTVKQLQALQONGSSVLSDEKSKRLNTILNTWSTIYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEMARANHYED 180
Db |||||
139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEMARANHYED 198
QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEETKPLYEHLHAYVRAKLMNAYSISYSP 240
Db |||||
199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEETKPLYEHLHAYVRAKLMNAYSISYSP 258
QY 241 IGCLPAHLGDMWGRFWNLVSLTVFPGQKPNIDVTDAMVDQAWDAQRIKFAEKEFFVSV 300
Db |||||
259 IGCLPAHLGDMWGRFWNLVSLTVFPGQKPNIDVTDAMVDQAWDAQRIKFAEKEFFVSV 318
QY 301 GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMGGH 360

```
Db 319 GLPNNTQGFWNSMLTDPGNVQAVCHPTAMDGLGKDFRILMCTKTMDDFLTAHHEMGH 378
Qy 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLAATPKHLKSLGSLSPDFQEDNETEIFN 420
Db 379 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLAATPKHLKSLGSLSPDFQEDNETEIFN 438
Qy 421 LLKQALTIIVGTLPTFTYMLEKWRWVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 480
Db 439 LLKQALTIIVGTLPTFTYMLEKWRWVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 498
Qy 481 DPASLFHVSNDYSFIRYTRTYLQFQOEALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 540
Db 499 DPASLFHVSNDYSFIRYTRTYLQFQOEALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 558
Qy 541 RLKSEPTWLTALENVVGAKNMVRPLLYNPEPLFTWLKQDNKNSFVGWSTDWSPY 595
Db 559 RLKSEPTWLTALENVVGAKNMVRPLLYNPEPLFTWLKQDNKNSFVGWSTDWSPY 613

RESULT 6
AAE20353
ID AAE20353 standard; protein; 805 AA.
XX
AC AAE20353;
XX
DT 18-JUN-2002 (first entry)
XX
DE Human ACE-2 full-length protein.
XX
KW Human; angiotensin converting enzyme-2; ACE-2; therapy; hypertension;
KW peptidyl dipeptidase A; blood pressure; hypotension; atherosclerosis;
KW myocardial infarction; heart failure; arrhythmia; renal failure; gene;
KW inflammation; fertility; enzyme; EC 3.4.15.1.
XX
OS Homo sapiens.

FH Key Location/Qualifiers
FT Peptide 1..18
FT Protein /label= Signal_peptide
FT Domain /label= Mature_ACE-2_protein
FT Domain 374..378
FT Domain /label= ZBD
FT Domain /note= "Zinc binding domain"
FT Domain 741..765
FT Domain /note= "Transmembrane domain; Hydrophobic region"
FT Domain 766..805
FT Domain /label= Cytoplasmic_domain
XX
US6194556-B1.
PN
XX
PD 27-FEB-2001.
XX
PF 11-DEC-1997; 97US-00989299.
XX
PR 11-DEC-1997; 97US-00989299.
XX
PA (WILL-) MILLENNIUM PHARM INC.
XX
PI Acton SL, Robison KE;
XX
XX WPI; 2001-210604/21.
XX
DR N-PSDB; AAD02758.
XX
XX
XX Novel genes encoding angiotensin converting enzyme-2 useful as antisense
XX or antigene agents for therapeutics, diagnostics and screening assays.
XX
PS Claim 33; Fig 1; 76pp; English.
XX
CC The present amino acid sequence is human angiotensin converting enzyme-2
XX (ACE-2), also referred as peptidyl dipeptidase A (EC 3.4.15.1). Nucleic
XX acid sequence encoding ACE-2 is useful as antisense or antigene agents
XX
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CC for sequence specific modulation of gene expression or in the analysis of
CC single base-pair mutations in the gene. Nucleic acid sequence encoding
CC ACE-2 is useful in therapeutics, diagnostics and in screening assays. ACE
CC -2 antagonist is used to treat hypertension or congestive heart failure
CC (CHF). ACE agonist is used to reduce the inflammation and pain resulting
CC from an insect sting or bite, which was accompanied by an injection of
CC bradykinin. Anti-ACE-2 antibodies are used to monitor ACE-2 protein
CC levels for determining the disease or condition associated with an
CC aberrant protein level
XX
SQ Sequence 805 AA;
Query Match 100.0%; Score 3231; DB 4; Length 805;
Best Local Similarity 100.0%; Pred. No. 6.1e-288;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 STIEEQAKTFDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKNSAFLEKQST 60
Db 19 STIEEQAKTFDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKNSAFLEKQST 78
Qy 61 LAQMYPLQEIQNLTVKLQALQONGSSVLSSEKSKRLNTILNTMTSTYSTCKVCPDNP 120
Db 79 LAQMYPLQEIQNLTVKLQALQONGSSVLSSEKSKRLNTILNTMTSTYSTCKVCPDNP 138
Qy 121 QECILLEPGLNEIMANSIDYNERLWAWESWSEVKQLRPLYEYVVLKNEVARAHYED 180
Db 139 QECILLEPGLNEIMANSIDYNERLWAWESWSEVKQLRPLYEYVVLKNEVARAHYED 198
Qy 181 YGDYWRGDEYVNGVDGYDSRGQLLEDVEHTFEEIKPLYEHLHAYVRAKLMNAYSISYP 240
Db 199 YGDYWRGDEYVNGVDGYDSRGQLLEDVEHTFEEIKPLYEHLHAYVRAKLMNAYSISYP 258
Qy 241 IGCLPAHLLGDMWGRFTNLYSLTVPPFGQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSU 300
Db 259 IGCLPAHLLGDMWGRFTNLYSLTVPPFGQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSU 318
Qy 301 GLPNNTQGFWNSMLTDPGNVQAVCHPTAMDGLGKDFRILMCTKTMDDFLTAHHEMGH 360
Db 319 GLPNNTQGFWNSMLTDPGNVQAVCHPTAMDGLGKDFRILMCTKTMDDFLTAHHEMGH 378
Qy 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLAATPKHLKSLGSLSPDFQEDNETEIFN 420
Db 379 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLAATPKHLKSLGSLSPDFQEDNETEIFN 438
Qy 421 LLKQALTIIVGTLPTFTYMLEKWRWVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 480
Db 439 LLKQALTIIVGTLPTFTYMLEKWRWVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 498
Qy 481 DPASLFHVSNDYSFIRYTRTYLQFQOEALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 540
Db 499 DPASLFHVSNDYSFIRYTRTYLQFQOEALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 558
Qy 541 RLKSEPTWLTALENVVGAKNMVRPLLYNPEPLFTWLKQDNKNSFVGWSTDWSPY 595
Db 559 RLKSEPTWLTALENVVGAKNMVRPLLYNPEPLFTWLKQDNKNSFVGWSTDWSPY 613

RESULT 7
AAE20353
ID AAE20353 standard; protein; 805 AA.
XX
AC AAE20353;
XX
DT 18-JUN-2002 (first entry)
XX
DE Human ACE-2 full-length protein.
XX
KW Human; angiotensin converting enzyme-2; ACE-2; therapy; hypertension;
KW peptidyl dipeptidase A; blood pressure; hypotension; atherosclerosis;
KW myocardial infarction; heart failure; arrhythmia; renal failure; gene;
KW inflammation; fertility; enzyme; EC 3.4.15.1.
XX
OS Homo sapiens.
```


XX The invention relates to an isolated nucleic acid from a human gene
 CC encoding aminopeptidase P (XPNEP2), bradykinin receptor B1 (BDKRB1),
 CC tachykinin receptor B1 (TACR1), C1 esterase inhibitor (C1NH), kallikrein
 CC 1 (KLK1), bradykinin receptor B2 (BDKRB2), angiotensin converting enzyme
 CC 2 (ACE2) or protease inhibitor 4 (PI4), comprising at least one
 CC polymorphic position. Also included are (1) a probe that hybridises to a
 CC polymorphic position as provided in the detailed summary of single
 CC nucleotide polymorphisms comprising additional 5' and 3' flanking genomic
 CC sequence; (2) analysing (M1) at least one nucleic acid sample comprising
 CC obtaining the sample from one or more individuals and determining the
 CC nucleic acid sequence at one or more polymorphic positions in a gene
 CC encoding a protein selected from the group above; (3) constructing (M2)
 CC haplotypes using the genes comprising grouping at least two nucleic acids
 CC ; (4) identifying (M3) an individual at risk of developing a disorder
 CC upon administration of an ACE inhibitor and/or vasopeptidase inhibitor
 CC using the polymorphic data; (5) a library of nucleic acids, each of which
 CC comprises one or more polymorphic positions within a gene encoding a
 CC human protein selected from the group above; and (6) genotyping (M4) an
 CC individual comprising obtaining a nucleic acid sample, determining the
 CC nucleotide present in at least one polymorphic position, and comparing at
 CC least one position with a known data set. The genes, (M1, M2, M3 and M4)
 CC and compositions are useful for detecting, diagnosing, treating,
 CC preventing various disorders such as angioedema and diseases which
 CC involve angiogenesis like haemangiomas, tumours, sarcomas, Crohn's
 CC disease, trachomas, and cardiovascular diseases like angina pectoris,
 CC hypertension, heart failure, myocardial infarction, ventricular
 CC hypertrophy, vascular diseases, aneurysm, embolism, thrombosis, coronary
 CC artery disease, arteriosclerosis and/or atherosclerosis, and
 CC hypersensitivity reactions, sepsis, autoimmune diseases, inflammatory
 CC arthritis, cancer, wounds, viral, bacterial or fungal infection, Chronic
 CC obstructive pulmonary disease (COPD) and enterocolitis (many other
 CC diseases and disorders are listed in the specification). The
 CC polynucleotides are also useful for chromosome identification. Antibodies
 CC against the proteins may be utilised for immunophenotyping of cell lines
 CC and biological samples. The present sequence represents one of the
 CC proteins listed above
 XX
 SX Sequence 805 AA;

Query Match 100.0%; Score 3231; DB 5; Length 805;
 Best Local Similarity 100.0%; Pred. No. 6.1e-288;
 Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAFTFLDKFNHEADFLYQSSLASWNTNTNTIENVQNMNAGDKSAFLKEQST 60
 DB 19 STIEEQAFTFLDKFNHEADFLYQSSLASWNTNTNTIENVQNMNAGDKSAFLKEQST 78
 QY 61 LAQMYPLOBIQNLTVKQLQALQONGSSVLSBDKSKRLNTILNTMTSTYTGKVCNPDNP 120
 DB 79 LAQMYPLOBIQNLTVKQLQALQONGSSVLSBDKSKRLNTILNTMTSTYTGKVCNPDNP 138
 QY 121 QECLLLEPGLINEIMANSLDYNRLWAWESWRSEVQKQLRPLYEEYVVLNWARANHVED 180
 DB 139 QECLLLEPGLINEIMANSLDYNRLWAWESWRSEVQKQLRPLYEEYVVLNWARANHVED 198
 QY 181 YGDYWRGDYEVANGDGYDSRGOLIEDVHTFEEIKPLYEHLHAYVRAKLMNAYSISF 240
 DB 199 YGDYWRGDYEVANGDGYDSRGOLIEDVHTFEEIKPLYEHLHAYVRAKLMNAYSISF 258
 QY 241 IGCPLPAHLIGDMWGRFWTNLYSLTVFPQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSFV 300
 DB 259 IGCPLPAHLIGDMWGRFWTNLYSLTVFPQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSFV 318
 QY 301 GLPNMTQGFWNSMLTRDGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMCH 360
 DB 319 GLPNMTQGFWNSMLTRDGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMCH 378
 QY 361 IQYDMAYAAQPFLLRNGANGEGHEAVEGTEIMSLSAATPKHLKSTGLSLSPDFQDNTEINF 420
 DB 379 IQYDMAYAAQPFLLRNGANGEGHEAVEGTEIMSLSAATPKHLKSTGLSLSPDFQDNTEINF 438
 QY 421 LLKQALTVTGLTLPFTYMLEKRWMMVKGEIPKQDMKKWEMKREIVGVVPEVPHDETTC 480

DB 439 LLKQALTVTGLTLPFTYMLEKRWMMVKGEIPKQDMKKWEMKREIVGVVPEVPHDETTC 498
 QY 481 DPASLPHVSNDSYSPRYRTYTRTYLQFQFBALCQAAKHEGHLKCDISNSTEAGQKLFNML 540
 DB 499 DPASLPHVSNDSYSPRYRTYTRTYLQFQFBALCQAAKHEGHLKCDISNSTEAGQKLFNML 558
 QY 541 RLKSKSEPTLALENVVGAQNMNVRPLLYFEPFLTWLKDQNKNSFVGWSTDSWSPY 595
 DB 559 RLKSKSEPTLALENVVGAQNMNVRPLLYFEPFLTWLKDQNKNSFVGWSTDSWSPY 613

RESULT 9
 ABG77023
 ID ABG77023 standard; protein; 805 AA.
 XX
 AC ABG77023;
 XX
 DT 05-NOV-2002 (first entry)
 XX
 DE Human angiotensin converting enzyme 2 variant #1.
 XX
 KW Aminopeptidase P; XPNEP2; bradykinin receptor B1; human; BDKRB1;
 KW tachykinin receptor B1; TACR1; C1 esterase inhibitor; C1NH; kallikrein 1;
 KW KLK1; bradykinin receptor B2; BDKRB2; gene therapy;
 KW angiotensin converting enzyme 2; ACE2; protease inhibitor 4; PI4;
 KW polymorphism; haemangioma; tumour; sarcoma; Crohn's disease; trachoma;
 KW cardiovascular disease; angina pectoris; hypertension; heart failure;
 KW myocardial infarction; ventricular hypertrophy; vascular disease;
 KW aneurysm; embolism; thrombosis; coronary artery disease; angioedema;
 KW arteriosclerosis; atherosclerosis; hypersensitivity; sepsis;
 KW autoimmune disease; inflammatory arthritis; cancer; wound;
 KW viral infection; bacterial infection; fungal infection; COPD;
 KW Chronic obstructive pulmonary disease; enterocolitis.
 XX
 OS Homo sapiens.
 XX
 XX WO200261131-A2.
 XX
 XX 08-AUG-2002.
 XX
 XX 03-DEC-2001; 2001WO-US047235.
 XX
 XX 04-DEC-2000; 2000US-0251015P.
 PR 23-JAN-2001; 2001US-0263678P.
 PR 02-MAR-2001; 2001US-0273037P.
 XX
 XX (BRIM) BRISTOL-MYERS SQUIBB CO.
 PA (TSUC/) TSUCHIHASHI Z.
 PA (HUIL/) HUI L.
 XX
 PI Tsuchihashi Z, Hui L, Zerba KB, Ma-Edmonds M, Perrone MH;
 Swanson BN, Powell JR;
 XX
 XX WPI; 2002-619265/66.
 DR N-PSDB; ABS60633.
 XX
 XX New isolated nucleic acid with at least one polymorphic position, useful
 PT for detecting, diagnosing and treating disorders such as angioedema,
 PT cancer, viral, bacterial or fungal infection, cardiovascular and
 PT autoimmune diseases.
 XX
 PS Disclosure; Fig 37; 977pp; English.
 XX
 CC The invention relates to an isolated nucleic acid from a human gene
 CC encoding aminopeptidase P (XPNEP2), bradykinin receptor B1 (BDKRB1),
 CC tachykinin receptor B1 (TACR1), C1 esterase inhibitor (C1NH), kallikrein
 CC 1 (KLK1), bradykinin receptor B2 (BDKRB2), angiotensin converting enzyme
 CC 2 (ACE2) or protease inhibitor 4 (PI4), comprising at least one
 CC polymorphic position. Also included are (1) a probe that hybridises to a
 CC polymorphic position as provided in the detailed summary of single
 CC nucleotide polymorphisms comprising additional 5' and 3' flanking genomic
 CC sequence; (2) analysing (M1) at least one nucleic acid sample comprising

obtaining the sample from one or more individuals and determining the nucleic acid sequence at one or more polymorphic positions in a gene encoding a protein selected from the group above; (3) constructing (M2) haplotypes using the genes comprising grouping at least two nucleic acids; (4) identifying (M3) an individual at risk of developing a disorder upon administration of an ACE inhibitor and/or vasopeptidase inhibitor using the polymorphic data; (5) a library of nucleic acids, each of which comprises one or more polymorphic positions within a gene encoding a human protein selected from the group above; and (6) genotyping (M4) an individual comprising obtaining a nucleic acid sample, determining the nucleotide present in at least one polymorphic position, and comparing at least one position with a known data set. The genes, (M1, M2, M3 and M4) and compositions are useful for detecting, diagnosing, treating, preventing various disorders such as angioedema and diseases which involve angiogenesis like haemangiomas, tumours, sarcomas, Crohn's disease, trachomas, and cardiovascular diseases like angina pectoris, hypertension, heart failure, myocardial infarction, ventricular artery disease, arteriosclerosis and/or atherosclerosis, and hypersensitivity reactions, sepsis, autoimmune diseases, inflammatory arthritis, cancer, wounds, viral, bacterial or fungal infection, Chronic obstructive pulmonary disease (COPD) and enterocolitis (many other diseases and disorders are listed in the specification). The polynucleotides are also useful for chromosome identification. Antibodies against the proteins may be utilised for immunophenotyping of cell lines and biological samples. The present sequence represents a polymorphic variant of one of the proteins listed above

Sequence 805 AA;

Query Match		100.0%;	Score 3231;	DB 5;	Length 805;
Best Local Similarity		100.0%;	Pred. No. 6.1e-289;		
Matches 595;		Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	STIEQAKTFLDKFNHEADLPYQSSLASWNTNTIENVQNMNAGDKWSAFLKEQST	60		
DB	19	STIEQAKTFLDKFNHEADLPYQSSLASWNTNTIENVQNMNAGDKWSAFLKEQST	78		
QY	61	LAQMYPLOEIQNLTVKQLQALQOQNGSSVLSDBKSKRLNTILNTWSTIYSTGKVCNPNP	120		
DB	79	LAQMYPLOEIQNLTVKQLQALQOQNGSSVLSDBKSKRLNTILNTWSTIYSTGKVCNPNP	138		
QY	121	QECLLLEPGLNEIMANSLDYNERLMAWESWRSEVKGQLRPLYEYVVLKNEANRANHYED	180		
DB	139	QECLLLEPGLNEIMANSLDYNERLMAWESWRSEVKGQLRPLYEYVVLKNEANRANHYED	198		
QY	181	YGDYWRGDEYVNGVDGYDSRGQIEDVHTFEEIKPLYEHLHAYVRKLMNAYPSYISP	240		
DB	199	YGDYWRGDEYVNGVDGYDSRGQIEDVHTFEEIKPLYEHLHAYVRKLMNAYPSYISP	258		
QY	241	IGCLPAHLIGDMGRFWTNLYSLTVPFQOKPNIDVTDAMVQAWDAQRIFKBAEKFVSV	300		
DB	259	IGCLPAHLIGDMGRFWTNLYSLTVPFQOKPNIDVTDAMVQAWDAQRIFKBAEKFVSV	318		
QY	301	GLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAHHEMGH	360		
DB	319	GLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAHHEMGH	378		
QY	361	IQYDMAYAAQPFLLRNGANEGPHEAVGEIMSLISAATPKHLKSGILLSPDFQEDNETEINF	420		
DB	379	IQYDMAYAAQPFLLRNGANEGPHEAVGEIMSLISAATPKHLKSGILLSPDFQEDNETEINF	438		
QY	421	LKQALTIIVGTLPFTYMLSEKRWMPVKGRIKPDQNMKKWEMKREIVGVVEVPVPHDETVC	480		
DB	439	LKQALTIIVGTLPFTYMLSEKRWMPVKGRIKPDQNMKKWEMKREIVGVVEVPVPHDETVC	498		
QY	481	DPASLPHVSNDSYFIRYRTLYLQFQFQALQAAKHGFLHKCDISNSTEAGQKLFNML	540		
DB	499	DPASLPHVSNDSYFIRYRTLYLQFQFQALQAAKHGFLHKCDISNSTEAGQKLFNML	558		
QY	541	RLGKSPWTLALENVVGAKNMVRPLLNYFEPFLTWLKDQNNKNSFVGWSTWSPY	595		
DB	559	RLGKSPWTLALENVVGAKNMVRPLLNYFEPFLTWLKDQNNKNSFVGWSTWSPY	613		

RESULT 10	
AAU99701	
ID	AAU99701 standard; protein; 805 AA.
XX	
AC	AAU99701;
XX	
DT	24-SEP-2002 (first entry)
XX	
DE	Human angiotensin converting enzyme-2 (ACE-2) protein.
XX	
KW	Human; angiotensin converting enzyme-2; ACE-2; body weight disorder;
KW	muscle mass; body fat; obesity; diabetes; atherosclerosis; weight loss;
KW	lipid metabolism; weight gain; anorexia; cachexia; bulimia; sepsis;
KW	familial partial lipodystrophy; hypercholesterolaemia; hyperlipidaemia;
KW	aberrant metabolic rate; heart failure; left ventricular hypertrophy;
KW	neurodegenerative disorder; peptide hormone; cytokine processing;
KW	myocardial infarction; cardiomyopathy; inflammatory bowel disease;
KW	systemic inflammation response syndrome; polytrauma; pain; stroke;
KW	bone destruction; rheumatoid arthritis; osteoarthritis; asthma;
KW	periodontal disease; dysmenorrhoea; premature labour; brain oedema;
KW	focal injury; diffuse axonal injury; reperfusion injury; scar formation;
KW	cerebral vasospasm; subarachnoid haemorrhage; allergic disorder;
KW	adult respiratory distress syndrome; wound healing; appetite;
KW	body mass index.
XX	
OS	Homo sapiens.
XX	
FH	Key
FT	Peptide
FT	1..18
FT	/label= Signal_peptide
FT	Protein
FT	19..805
FT	/label= Mature_human_ACE_2_protein
XX	
WO	200239997-A2.
XX	
PD	23-MAY-2002.
XX	
PF	31-OCT-2001; 2001WO-US045703.
XX	
PR	01-NOV-2000; 2000US-00704216.
PR	29-MAY-2001; 2001US-00870382.
PR	19-OCT-2001; 2001US-0371741P.
XX	
PA	(MILL-) MILLENNIUM PHARM INC.
XX	
PI	Acton SL, Ocain TD, Gould AE, Dales NA, Guan B, Brown JA;
PI	Patane M, Kadambi Vu, Solomon M, Stricker-Krongrad A;
XX	
DR	WPI: 2002-547572/58.
DR	N-PSDB; ABK87623.
XX	
PT	Treating body weight disorder and increasing muscle mass comprises
PT	administering angiotensin converting enzyme-2 modulating compound.
XX	
PS	Example 5; Page 387-390; 395pp; English.
XX	
CC	The present invention describes a new method of treating a body weight
CC	disorder, increasing muscle mass and decreasing body fat by
CC	administration of angiotensin converting enzyme (ACE)-2 modulating
CC	compound. The invention can be used for treating body weight disorders,
CC	particularly obesity of at least grade 1, diabetes, atherosclerosis and a
CC	state associated with lipid metabolism. The method is used for treating
CC	rapid weight loss, rapid weight gain, anorexia, cachexia, bulimia,
CC	generalised partial lipodystrophy, familial partial lipodystrophy,
CC	hypercholesterolaemia, hyperlipidaemia, an aberrant metabolic rate,
CC	congestive heart failure, chronic heart failure, left ventricular
CC	hypertrophy, acute heart failure, neurodegenerative disorders (e.g.
CC	Alzheimer's disease, Parkinson's disease and Huntington's disease),
CC	diseases associated with peptide hormones or cytokine processing,
CC	myocardial infarction, cardiomyopathy, systemic inflammation response
CC	syndrome, sepsis, polytrauma, inflammatory bowel disease, acute and

CC	chronic pain, bone destruction in rheumatoid arthritis and osteoarthritis
CC	and periodontal disease, dysmenorrhea, premature labour, brain oedema
CC	following focal injury, diffuse axonal injury, stroke, reperfusion
CC	injury, cerebral vasospasm after subarachnoid haemorrhage, allergic
CC	disorders including asthma, adult respiratory distress syndrome, wound
CC	healing and scar formation. The invention decreases the appetite,
CC	increases muscle mass and decreases body fat of subject having body mass
CC	index of greater than 23 (preferably 24.9)kg/m ² . The present amino acid
XX	sequence represents the human ACE-2 protein of the invention
XX	
SQ	Sequence 805 AA;
Query Match 100.0%; Score 3231; DB 5; Length 805;	
Best Local Similarity 100.0%; Pred. No. 6.1e-288;	
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
QY	1 STIEQAKTFLDKFNHEADLFYQSSLASWNTNTIENVMNAGDKWSAFLKEQST 60
Db	19 STIEQAKTFLDKFNHEADLFYQSSLASWNTNTIENVMNAGDKWSAFLKEQST 78
QY	61 LAQMPLOEIQNLTVKQLQALQONGSSVLSDEKSKRLNTILNTMTSTIYTGKVCNPDNP 120
Db	79 LAQMPLOEIQNLTVKQLQALQONGSSVLSDEKSKRLNTILNTMTSTIYTGKVCNPDNP 138
QY	121 QECLLLEPGLNEIMANSLDYNRLWAWESVSGKQLRPLYEYVVLKNEVARANHYED 180
Db	139 QECLLLEPGLNEIMANSLDYNRLWAWESVSGKQLRPLYEYVVLKNEVARANHYED 198
QY	181 YGDYWRGDYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYSISYP 240
Db	199 YGDYWRGDYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYSISYP 258
QY	241 IGCPLPAHLGDMWGRFWTNLYSLTVPFQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 300
Db	259 IGCPLPAHLGDMWGRFWTNLYSLTVPFQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 318
QY	301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRLMCTKVTMDDELTAHHEMCH 360
Db	319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRLMCTKVTMDDELTAHHEMCH 378
QY	361 IOYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSLGILLSPDFQEDNETINF 420
Db	379 IOYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSLGILLSPDFQEDNETINF 438
QY	421 LKQALTIIVGTLPFTYMLBKRWMPKGBIPKQDMKKWEMKREITGVVPEVPHDETYC 480
Db	439 LKQALTIIVGTLPFTYMLBKRWMPKGBIPKQDMKKWEMKREITGVVPEVPHDETYC 498
QY	481 DPASLPHVNSDYSFIRYTRTLYQFOFQALCOAKHEGFLHKCDISNSTEAGOKLFNNML 540
Db	499 DPASLPHVNSDYSFIRYTRTLYQFOFQALCOAKHEGFLHKCDISNSTEAGOKLFNNML 558
QY	541 RLKGSPPWTLALENVVGAKMNVRLPLNTYFPEPLFTWLKDQKNKSFVGVSTWDSPY 595
Db	559 RLKGSPPWTLALENVVGAKMNVRLPLNTYFPEPLFTWLKDQKNKSFVGVSTWDSPY 613
RESULT 11	
ID	ABU07731
XX	ABU07731 standard; protein; 805 AA.
AC	ABU07731;
XX	
XX	27-MAY-2003 (first entry)
DT	
XX	Human zinc metalloproteinase Zace2.
DE	
XX	Human; enzyme; Zace2; zinc metalloproteinase; ulcerative colitis;
KW	inflammation; inflammatory bowel disease; arthritis; enterocolitis;
KW	Crohn's disease; gene therapy; transgenic.
OS	
XX	Homo sapiens.
XX	

PH	Key	Location/Qualifiers
FT	Region	371..380
FT	Region	/label= Expanded_zinc_binding_region
FT	Region	374..378
FT	Region	/label= Zinc-binding_motif
FT	Domain	739..761
FT	Domain	/label= Transmembrane_domain
XX		
PN	US2002177211-A1.	
XX		
PD	28-NOV-2002.	
XX		
XX	16-OCT-2001; 2001US-00978385.	
PF		
XX	13-MAY-1999; 99US-0133952P.	
PR	27-AUG-1999; 99US-0151181P.	
PR	03-MAY-2000; 2000US-00563516.	
XX		
XX	(ZYMO) ZYMOGENETICS INC.	
XX		
PI	Piddington CS, Petrie C, Shoemaker KE, Bishop PD;	
XX	WPI; 2003-328489/31.	
DR	N-PSDB; ABX93333.	
XX		
XX	Isolated human or murine Zace2 polypeptide useful for reducing	
PT	inflammation in conditions such as inflammatory bowel disease, arthritis,	
PT	enterocolitis, ulcerative colitis and Crohn's disease.	
XX		
PS	Claim 1; Page 39-41; 57pp; English.	
XX		
CC	The invention relates to an isolated polypeptide, comprising fully	
CC	defined human Zace2, murine Zace-5, or murine Zace2-10 polypeptide. An	
CC	expression vector containing Zace2 polynucleotide is useful for producing	
CC	Zace2 protein. The polynucleotide is useful as a diagnostic probe for	
CC	detecting a product of Zace2 gene expression in a biological sample. The	
CC	polypeptide is also useful for decreasing inflammation associated with a	
CC	condition such as inflammatory bowel disease, arthritis or enterocolitis.	
CC	The polypeptide is also useful for treating Crohn's disease and	
CC	ulcerative colitis. The polypeptide is useful for producing labelled	
CC	angiotensin II, for identifying modulators of zinc protease activity and	
CC	for identifying angiotensin converting enzyme (ACE) inhibitors. The	
CC	polynucleotide is useful in gene therapy techniques to treat the above	
CC	mentioned disorders. The polynucleotide is also useful for determining	
CC	whether a subject's chromosome contains a mutation in the Zace2 gene. The	
CC	present sequence represents the amino acid sequence of human zinc	
CC	metalloproteinase Zace2	
XX		
SQ	Sequence 805 AA;	
Query Match 100.0%; Score 3231; DB 6; Length 805;		
Best Local Similarity 100.0%; Pred. No. 6.1e-288;		
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
QY	1	STIEQAKTFLDKFNHEADLFYQSSLASWNTNTIENVMNAGDKWSAFLKEQST 60
Db	19	STIEQAKTFLDKFNHEADLFYQSSLASWNTNTIENVMNAGDKWSAFLKEQST 78
QY	61	LAQMPLOEIQNLTVKQLQALQONGSSVLSDEKSKRLNTILNTMTSTIYTGKVCNPDNP 120
Db	79	LAQMPLOEIQNLTVKQLQALQONGSSVLSDEKSKRLNTILNTMTSTIYTGKVCNPDNP 138
QY	121	QECLLLEPGLNEIMANSLDYNRLWAWESVSGKQLRPLYEYVVLKNEVARANHYED 180
Db	139	QECLLLEPGLNEIMANSLDYNRLWAWESVSGKQLRPLYEYVVLKNEVARANHYED 198
QY	181	YGDYWRGDYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYSISYP 240
Db	199	YGDYWRGDYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYSISYP 258
QY	241	IGCPLPAHLGDMWGRFWTNLYSLTVPFQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 300
Db	259	IGCPLPAHLGDMWGRFWTNLYSLTVPFQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 318

QY 301 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMWH 360
DB 319 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMWH 378
QY 361 IQYDMAYAAQPFLLRNGANEGFHEAVGETIMSLSAATPKHLKXIGLLSPDFQEDNETEINF 420
DB 379 IQYDMAYAAQPFLLRNGANEGFHEAVGETIMSLSAATPKHLKXIGLLSPDFQEDNETEINF 438
QY 421 LKQALTIIVGTLPFTYMLKRWMMVFKGEIPKQOMKWKWEMKREIVGVVEVPVHDETYC 480
DB 439 LKQALTIIVGTLPFTYMLKRWMMVFKGEIPKQOMKWKWEMKREIVGVVEVPVHDETYC 498
QY 481 DPASLPHVNSDYSFIRYTRTYLQFOFQALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 540
DB 499 DPASLPHVNSDYSFIRYTRTYLQFOFQALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 558
QY 541 RLKSEPTWLALENVVGAKNMVRPLNYPFELFTWLKQDNKNSFVGWSTDSWSPY 595
DB 559 RLKSEPTWLALENVVGAKNMVRPLNYPFELFTWLKQDNKNSFVGWSTDSWSPY 613
RESULT 12
ID ADA03344
XX ADA03344 standard; protein; 805 AA.
AC ADA03344;
XX
DT 06-NOV-2003 (first entry)
XX Human angiotensin converting enzyme 2-like protein.
XX
KW hypotensive; cardiant; cerebroprotective; antiatherosclerotic; analgesic;
KW antiinflammatory; nephrotropic; hypertensive; vasotropic; cyostatic;
KW antiasthmatic; antiallergic; neuroprotective; antiparkinsonian;
KW nootropic; antirheumatic; antiarthritic; antigout; tranquilizer;
KW vulnary; antidiabetic; dermatological; immunosuppressive; hepatotropic;
KW anti-HIV; antibacterial; angiotensin converting enzyme; ACE-2;
KW angiotensin converting enzyme; ACE-2; hypertension;
KW congestive heart failure; stroke; left ventricular failure;
KW atherosclerotic heart disease; stenosis; pain; inflammatory reaction;
KW histamine; vasoconstriction; epitope; aldosterone; cell proliferation;
KW renal disorder; acute glomerulonephritis; immunophenotyping;
KW cardiac myocyte; Bowman's capsule; hypotensin; ischemia; asthma; allergy;
KW multiple sclerosis; cancer; Parkinson's disease; Alzheimer's disease;
KW rheumatoid arthritis; gout; trauma; dermatitis; diabetes mellitus;
KW Sjogren's syndrome; Addison's disease; hepatitis; Crohn's disease;
KW sarcoidosis; AIDS; sepsis.
XX
OS Homo sapiens.
XX
PN WO200298448-A1.
XX
XX 12-DEC-2002.
XX
XX 03-JUN-2002; 2002WO-US017199.
XX
XX 04-JUN-2001; 2001US-0294976P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
PI Parry TJ, Rosen CA, Albert VR, Sanyal I, Huang L, Wescott CR;
PI Sekut L;
XX
XX WPI; 2003-140552/13.
DR N-PSDB; ADA03343.
XX
XX Novel angiotensin converting enzyme-2 binding polypeptide useful for
PT treating, preventing or ameliorating hypertension, congestive heart
PT failure, stroke, left ventricular failure and atherosclerotic heart
PT disease.
XX
PS Disclosure; Page 239-241; 246pp; English.

XX The invention relates to novel isolated angiotensin converting enzyme
CC (ACE)-2 binding polypeptides (I), which are useful for treating.
CC preventing or ameliorating hypertension, congestive heart failure, an
CC stroke, left ventricular failure and atherosclerotic heart disease in an
CC animal. The peptides are useful for detecting, isolating, or purifying
CC ACE-2 proteins or ACE-2 like polypeptides in solutions, mixtures, or
CC biological samples; for inhibiting or reducing stenosis, pain,
CC inflammatory reactions, abnormal histamine release, vasoconstriction,
CC diseases or disorders related to vasoconstriction, and diseases and/or
CC disorders associated with aberrant action of ACE-2; to detect, isolate,
CC or remove ACE-2 target proteins in solutions, and also to identify
CC epitopes of ACE-2; to detect, diagnose, prognose, or monitor
CC cardiovascular diseases, and disorders associated with aberrant
CC aldosterone activity, or cell proliferation; for preventing and treating
CC renal disorders, e.g., acute glomerulonephritis, and diseases associated
CC with it; to assay protein levels in a biological sample, for
CC immunophenotyping of cell lines and biological samples by their ACE-2
CC expression, and for identifying cells, such as cardiac myocytes,
CC endothelial and epithelial cells of Bowman's capsule. The peptides are
CC especially useful for treating, preventing, or ameliorating diseases or
CC disorders associated with hypotensin, ischemia, asthma, allergy, multiple
CC sclerosis, cancers, Parkinson's and Alzheimer's diseases, rheumatoid
CC arthritis, gout, trauma, dermatitis, diabetes mellitus, Sjogren's
CC syndrome, Addison's disease, chronic active hepatitis, Crohn's disease,
CC sarcoidosis, AIDS, and sepsis. In an example of the invention, ACE-2
CC binding peptides were isolated from a number of peptide display
CC libraries. Evaluation of the peptide sequences revealed a series of
CC peptide families. This sequence represents a human angiotensin converting
CC enzyme 2-like protein.
XX
SQ Sequence 805 AA;

Query Match 100.0%; Score 3231; DB 6; Length 805;
Best Local Similarity 100.0%; Pred. No. 6.1e-288;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 STIEQAKTFLDKFNHEADLFYQSSLASWNYNTNTEENVQNMNAGDKWSAFLEKEST 60
DB 19 STIEQAKTFLDKFNHEADLFYQSSLASWNYNTNTEENVQNMNAGDKWSAFLEKEST 78
QY 61 LAQMPYPLQEIQNLTKVLQALQALQNGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 120
DB 79 LAQMPYPLQEIQNLTKVLQALQALQNGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 138
QY 121 QECILLEFGLNEIMANSLDYNERLWAWESWSEVKQLRPLYEEYVVLKNEWARAHYED 180
DB 139 QECILLEFGLNEIMANSLDYNERLWAWESWSEVKQLRPLYEEYVVLKNEWARAHYED 198
QY 181 YGDYWRGDYVNGVDYDSRGOLIEDVEHTPEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
DB 199 YGDYWRGDYVNGVDYDSRGOLIEDVEHTPEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
QY 241 IGCLPAHLGDMWGRFTNLYSLTVPFGOKNIDVTDAMVQAWDAQRIKFAEKEFFVSV 300
DB 259 IGCLPAHLGDMWGRFTNLYSLTVPFGOKNIDVTDAMVQAWDAQRIKFAEKEFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMWH 360
DB 319 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVMTDDFLTAAHEMWH 378
QY 361 IQYDMAYAAQPFLLRNGANEGFHEAVGETIMSLSAATPKHLKXIGLLSPDFQEDNETEINF 420
DB 379 IQYDMAYAAQPFLLRNGANEGFHEAVGETIMSLSAATPKHLKXIGLLSPDFQEDNETEINF 438
QY 421 LKQALTIIVGTLPFTYMLKRWMMVFKGEIPKQOMKWKWEMKREIVGVVEVPVHDETYC 480
DB 439 LKQALTIIVGTLPFTYMLKRWMMVFKGEIPKQOMKWKWEMKREIVGVVEVPVHDETYC 498
QY 481 DPASLPHVNSDYSFIRYTRTYLQFOFQALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 540
DB 499 DPASLPHVNSDYSFIRYTRTYLQFOFQALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 558

QY 541 RLKSEPTLALENVVGAKNMVRPLINYPPEPLFTWLKDQNKSPVGNSTDMSPY 595
 DB 559 RLKSEPTLALENVVGAKNMVRPLINYPPEPLFTWLKDQNKSPVGNSTDMSPY 613

RESULT 13

ABR56712
 ID ABR56712 standard; protein; 805 AA.

XX AC ABR56712;
 XX 30-JUL-2003 (first entry)
 XX Human ACE-2 protein SEQ ID NO:142.
 XX Human; angiotensin converting enzyme 2; ACE-2 binding; ACE-2;
 KW vasoconstriction; low blood pressure; angiotensin II; angiotensin;
 KW hypertensive; vasotropic; vaccine; hypotension; shock; syncope.

XX Homo sapiens.

PN WO200298906-A1.

XX 12-DEC-2002.

PF 03-JUN-2002; 2002WO-US017213.

XX 04-JUN-2001; 2001US-0295004P.

XX (HUMA-) HUMAN GENOME SCI INC.

XX Parry TJ;

DR WPI; 2003-140586/13.

DR N-PSDB; ACC79022.

XX Increasing vasoconstriction or ameliorating a disorder resulting from low
 PT blood pressure, e.g. hypotension, shock or syncope, comprises
 PT administering an angiotensin II in combination with angiotensin 1-9 to an
 PT individual.

PS Disclosure; Page 227-229; 237pp; English.

XX The present invention describes a method for increasing vasoconstriction
 CC or ameliorating a disorder resulting from low blood pressure, which
 CC comprises administering to an individual an amount of angiotensin II in
 CC combination with angiotensin 1-9. Angiotensin has hypertensive and
 CC vasotropic activities, and can be used in vaccines. The method is useful
 CC for increasing vasoconstriction or ameliorating a disorder resulting from
 CC low blood pressure, such as hypotension, shock or syncope. ABR56563 to
 CC ABR56708 represent angiotensin converting enzyme 2 (ACE-2) binding
 CC peptides, ABR56709 to ABR56725 and ACC79021 to ACC79025 represent
 CC sequences used in the exemplification of the present invention. Human ACE
 CC -2 is located to chromosome X, more specifically to Xp22. N.B. ABR56563
 CC to ABR56572 represent SEQ ID NO:1 to 10 and should be the same as
 CC ABR56573 to ABR56582, but the Z's given at the beginning and end of the
 CC peptides in the disclosure have been expanded to Glx in the Sequence
 CC Listing and in this case the Z's do not represent Gln or Glu (see pages 4
 CC to 7). SEQ ID NO:40 to 136 in the Sequence Listing (see also pages 7 to
 CC 10) have been specified as SEQ ID NO:20 to 116 in Example 1 (see pages
 CC 174 to 177)

XX Sequence 805 AA;

Query Match 100.0%; Score 3231; DB 6; Length 805;
 Best Local Similarity 100.0%; Pred. No. 6.1e-286; Indels 0; Gaps 0;
 Matches 595; Conservative 0; Mismatches 0;

QY 1 STIEEQATFLDKFNHEADLFIQSSLSASWNTNTNTNTEENVQNMNAGDKWSAFLKEQST 60
 DB 19 STIEEQATFLDKFNHEADLFIQSSLSASWNTNTNTNTEENVQNMNAGDKWSAFLKEQST 78

QY 61 LAQMPLOEQIQLTVKQLQALQQNGSSVLSDKSKRLNTILNTWSTIYSTGKVCNPDNP 120

DB 79 LAQMPLOEQIQLTVKQLQALQQNGSSVLSDKSKRLNTILNTWSTIYSTGKVCNPDNP 138
 QY 121 QECLLLEPGLNEIMANSLDYNERLWAWBSRSEVGKQLRPPLYEEYVVLKXNEMARAHYED 180
 DB 139 QECLLLEPGLNEIMANSLDYNERLWAWBSRSEVGKQLRPPLYEEYVVLKXNEMARAHYED 198
 QY 181 YGDYWRGDIYEVNGVDGYDYSRQQLIEDVEHTPEEIKPLYEHLHAYYRAKLMNAYPSYISP 240
 DB 199 YGDYWRGDIYEVNGVDGYDYSRQQLIEDVEHTPEEIKPLYEHLHAYYRAKLMNAYPSYISP 258
 QY 241 IGCLPAHLGDMWGRFTNLXSLTVPFGQKPNIDVTDAWVDAQDAQRIKFAEKFFVSV 300
 DB 259 IGCLPAHLGDMWGRFTNLXSLTVPFGQKPNIDVTDAWVDAQDAQRIKFAEKFFVSV 318
 QY 301 GLPNNTQGFWENSLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVTMDDFLTAHHEMGH 360
 DB 319 GLPNNTQGFWENSLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVTMDDFLTAHHEMGH 378
 QY 361 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSGILLSPDFQSDNETEINF 420
 DB 379 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSGILLSPDFQSDNETEINF 438
 QY 421 LLKQALTTIVGTLPTFTYMLEKRWMMVFKGEIPKQOMKMKWEMKREITGVVSPVPHDETVC 480
 DB 439 LLKQALTTIVGTLPTFTYMLEKRWMMVFKGEIPKQOMKMKWEMKREITGVVSPVPHDETVC 498
 QY 481 DPASLFHVSNDYSFIRYTRTLYQFQFALCQAAKHGEPHLKCDISNSTEAGQKLFNNML 540
 DB 499 DPASLFHVSNDYSFIRYTRTLYQFQFALCQAAKHGEPHLKCDISNSTEAGQKLFNNML 558
 QY 541 RLKSEPTLALENVVGAKNMVRPLINYPPEPLFTWLKDQNKSPVGNSTDMSPY 595
 DB 559 RLKSEPTLALENVVGAKNMVRPLINYPPEPLFTWLKDQNKSPVGNSTDMSPY 613

RESULT 14

ADL95395
 ID ADL95395 standard; protein; 805 AA.

XX AC ADL95395;
 XX 20-MAY-2004 (first entry)
 XX Human angiotensin converting enzyme-2 (ACE-2).
 DE Human angiotensin converting enzyme-2 (ACE-2).
 KW bioactivity; angiotensin converting enzyme-2; ACE-2; human; enzyme;
 KW carboxypeptidase.

XX Homo sapiens.

XX US6610497-B1.

XX 26-AUG-2003.

XX 29-SEP-1999; 99US-00407427.

XX 11-DEC-1997; 97US-00989299.

XX 30-SEP-1998; 98US-00163648.

XX (MILL-) MILLENNIUM PHARM INC.

XX Acton SL, Robison KE, Hsieh FY;

XX WPI; 2003-895335/82.

XX N-PSDB; ADL95394, ADL95396.

XX Identification of compound that modulates bioactivity of angiotensin
 PT converting enzymes-2 polypeptide, by detecting modulation of the
 PT bioactivity of polypeptide that is contacted with test compound as
 PT compared to control.

XX Claim 6; SEQ ID NO 2; 91pp; English.

XX The invention describes a compound that modulates bioactivity of an
CC angiotensin converting enzyme-2 (ACE-2) polypeptide. The compound is
CC identified by contacting an ACE-2 polypeptide with a test compound under
CC conditions for modulation of the bioactivity of the polypeptide; and
CC detecting modulation of the bioactivity of the polypeptide by the test
CC compound as compared to a control. Also described is a method for
CC modulating the bioactivity of an ACE-2 polypeptide by contacting the ACE-
CC 2 polypeptide with a compound that has been identified. The method is
CC useful for identifying a compound that modulates the bioactivity of
CC angiotensin converting enzyme-2 peptides. The inventive method identifies
CC other potential substrates of ACE-2 polypeptides and the product of the
CC enzymatic reaction. The comparison of the mass spectra of the test
CC compound with that of the reaction mixture after incubation indicates
CC whether the test compound was converted into a new compound, in which
CC case the test compound is a substrate of the ACE-2 polypeptide. This is
CC the amino acid sequence of human angiotensin converting enzyme-2 (ACE-2).
XX
SQ Sequence 805 AA;
Query Match 100.0%; Score 3231; DB 7; Length 805;
Best Local Similarity 100.0%; Pred. No. 6.1e-288; Indels 0; Gaps 0;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 STIEQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWSAFLKEQST 60
Db |||||
QY 19 STIEQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWSAFLKEQST 78
Db |||||
QY 61 LAQMPLOEIQNLTKVQLQALQONGSSVLSDEKSKRLNTILNTMTSTYTGKVCNPNP 120
Db |||||
QY 79 LAQMPLOEIQNLTKVQLQALQONGSSVLSDEKSKRLNTILNTMTSTYTGKVCNPNP 138
Db |||||
QY 121 QECILLEPGLNEIMANSLDYNRLWAWESRSEVGKQLRPLYEEYVVLKNEWARANHYED 180
Db |||||
QY 139 QECILLEPGLNEIMANSLDYNRLWAWESRSEVGKQLRPLYEEYVVLKNEWARANHYED 198
Db |||||
QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
Db |||||
QY 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
Db |||||
QY 241 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDMVDQAWDAQRIKFAEKFFVSV 300
Db |||||
QY 259 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDMVDQAWDAQRIKFAEKFFVSV 318
Db |||||
QY 301 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVMDDELTAHHEMGH 360
Db |||||
QY 319 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVMDDELTAHHEMGH 378
Db |||||
QY 361 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSGILLSPDFQEDNTEINF 420
Db |||||
QY 379 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSGILLSPDFQEDNTEINF 438
Db |||||
QY 421 LKQALTIIVGTLPTFTYMLEKRWMPFKGBI PKDQNMKKWEMKREIVGVVEVPVPHDETYC 480
Db |||||
QY 439 LKQALTIIVGTLPTFTYMLEKRWMPFKGBI PKDQNMKKWEMKREIVGVVEVPVPHDETYC 498
Db |||||
QY 481 DPASLPHVSNDSYFRYYRTLYQFQFALCOAKHGEGLPKHCDISNSTEAGQKLFNNL 540
Db |||||
QY 499 DPASLPHVSNDSYFRYYRTLYQFQFALCOAKHGEGLPKHCDISNSTEAGQKLFNNL 558
Db |||||
QY 541 RLKSKSPWTLALENVVGAKNMVRPLNLYFEPLFTWLKQDNKNSFVGWSTWDSPY 595
Db |||||
QY 559 RLKSKSPWTLALENVVGAKNMVRPLNLYFEPLFTWLKQDNKNSFVGWSTWDSPY 613
Db |||||

RESULT 15

ADL95494

ID ADL95494 standard; protein; 805 AA.

XX ADL95494;

AC ADL95494;

XX 20-MAY-2004 (first entry)

DT 20-MAY-2004 (first entry)

DE Human angiotensin converting enzyme-2 (ACE-2) N720D.
XX bioactivity; angiotensin converting enzyme-2; ACE-2; human; enzyme;
KW mutant; mutain.
XX Homo sapiens.
OS Synthetic.
XX Key Location/Qualifiers
FH Misc-difference 720 /note= "Wild type Asn substituted by Asp"
FT US6610497-B1.
XX 26-AUG-2003.
PD 29-SEP-1999; 99US-00407427.
XX 11-DEC-1997; 97US-00989299.
PR 30-SEP-1998; 98US-00163648.
PA (MILL-) MILLENNIUM PHARM INC.
XX Acton SL, Robison KE, Hsieh FY;
PI WPI; 2003-895335/82.
XX N-PSDB; ADL95394.
DR Identification of compound that modulates bioactivity of angiotensin
XX converting enzymes-2 polypeptide, by detecting modulation of the
PT bioactivity of polypeptide that is contacted with test compound as
PT compared to control.
XX Disclosure; SEQ ID NO 100; 91pp; English.
XX The invention describes a compound that modulates bioactivity of an
CC angiotensin converting enzyme-2 (ACE-2) polypeptide. The compound is
CC identified by contacting an ACE-2 polypeptide with a test compound under
CC conditions for modulation of the bioactivity of the polypeptide; and
CC detecting modulation of the bioactivity of the polypeptide by the test
CC compound as compared to a control. Also described is a method for
CC modulating the bioactivity of an ACE-2 polypeptide by contacting the ACE-
CC 2 polypeptide with a compound that has been identified. The method is
CC useful for identifying a compound that modulates the bioactivity of
CC angiotensin converting enzyme-2 peptides. The inventive method identifies
CC other potential substrates of ACE-2 polypeptides and the product of the
CC enzymatic reaction. The comparison of the mass spectra of the test
CC compound with that of the reaction mixture after incubation indicates
CC whether the test compound was converted into a new compound, in which
CC case the test compound is a substrate of the ACE-2 polypeptide. This is
CC the amino acid sequence of human angiotensin converting enzyme-2 (ACE-2)
CC N720D mutant. Note: This sequence does not appear in the printed
CC specification but has been created by the indexer using information given
CC in the invention.
XX
SQ Sequence 805 AA;
Query Match 100.0%; Score 3231; DB 7; Length 805;
Best Local Similarity 100.0%; Pred. No. 6.1e-288; Indels 0; Gaps 0;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 STIEQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWSAFLKEQST 60
Db |||||
QY 19 STIEQAKTFLDKFNHEADLFYQSSLASWNTNTNITEENVQNMNAGDKWSAFLKEQST 78
Db |||||
QY 61 LAQMPLOEIQNLTKVQLQALQONGSSVLSDEKSKRLNTILNTMTSTYTGKVCNPNP 120
Db |||||
QY 79 LAQMPLOEIQNLTKVQLQALQONGSSVLSDEKSKRLNTILNTMTSTYTGKVCNPNP 138
Db |||||
QY 121 QECILLEPGLNEIMANSLDYNRLWAWESRSEVGKQLRPLYEEYVVLKNEWARANHYED 180
Db |||||
QY 139 QECILLEPGLNEIMANSLDYNRLWAWESRSEVGKQLRPLYEEYVVLKNEWARANHYED 198
Db |||||

Qy 181 YGDYWRGDYEVNGVDGYDSRGOLIRDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
199 YGDYWRGDYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
Qy 241 IGCLPAHLLGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRIFKEAEKEFFVSU 300
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
259 IGCLPAHLLGDMWGRFWTNLYSLTVPGQKPNIDVTDAMVDQAWDAQRIFKEAEKEFFVSU 318
Qy 301 GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGDFRILMCTKVMTDDFLTAHHEMGGH 360
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
319 GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGDFRILMCTKVMTDDFLTAHHEMGGH 378
Qy 361 IOYDMAYAAQPFLLRNCANEGFHEAYGEIMSLSAATPKHLKSTGLLSPDFQEDNETEINF 420
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
379 IOYDMAYAAQPFLLRNCANEGFHEAYGEIMSLSAATPKHLKSTGLLSPDFQEDNETEINF 438
Qy 421 LLKQALTIUGTLPTTYMLEKWRWVPKGEIPKQWKKWEMKREIVGVVEPVPHDETYC 480
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
439 LLKQALTIUGTLPTTYMLEKWRWVPKGEIPKQWKKWEMKREIVGVVEPVPHDETYC 498
Qy 481 DPASLFHVSNDYSFIRYTYTTLTYQOFQEQALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 540
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
499 DPASLFHVSNDYSFIRYTYTTLTYQOFQEQALCOAAKHGEGPLHKCDISNSTEAGQKLFNML 558
Qy 541 RLCKSEBPTLALENVVCAKNMVRPLLNYPFLTWLKOQNKNSFVGWSTDWSPY 595
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
559 RLCKSEBPTLALENVVCAKNMVRPLLNYPFLTWLKOQNKNSFVGWSTDWSPY 613

Search completed: March 28, 2006, 11:09:51
Job time : 137.885 secs

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Result No.	Score	Query Match	Length	DB	ID	Description
1	3231	100.0	804	2	T14762	hypothetical prote
2	1335	41.3	732	1	S05238	peptidyl-dipeptida
3	1335	41.3	1306	1	A33759	peptidyl-dipeptida
4	1334	41.3	732	1	A33655	peptidyl-dipeptida
5	1334	41.3	1312	1	A34171	peptidyl-dipeptida
6	1310	40.5	1313	1	JC2038	peptidyl-dipeptida
7	1307	40.5	1193	2	JC2489	peptidyl-dipeptida
8	1281	39.6	737	1	A34402	peptidyl-dipeptida
9	1281	39.6	1309	1	S35484	peptidyl-dipeptida
10	1054.5	32.6	611	2	S65472	peptidyl-dipeptida
11	1027.5	31.8	630	2	JC5374	angiotensin-conver
12	1022	31.6	615	2	A57533	peptidyl-dipeptida
13	635.5	19.7	907	2	T15792	hypothetical prote
14	157	4.9	532	2	C83696	hypothetical prote
15	154	4.8	502	2	AF1310	probable thermosta
16	147	4.5	502	2	AE1682	probable thermosta
17	139.5	4.3	987	2	AI2011	peptide synthetase
18	139	4.3	608	2	B82938	zinc metalloprotei
19	136	4.2	611	2	D82881	zinc metalloprotei
20	135	4.2	501	2	D69943	carboxypeptidase h
21	124	3.8	538	2	E2561	probable thermosta
22	121	3.7	607	2	AB3511	oligoendopeptidase
23	121	3.7	1034	2	T30574	beta-galactosidase
24	118	3.7	627	1	S40048	1,4-alpha-glucan b
25	117.5	3.6	987	2	I48373	G-utrophin - mouse
26	115.5	3.6	611	2	A75573	probable oligoendo
27	114	3.5	685	2	F75370	oligoendopeptidase A
28	113.5	3.5	3655	2	T38084	TRAP-like protein
29	113	3.5	772	2	AI0968	probable glycosyl

2

2

2

Qy	1	STTIEQAKTFLDKFNHEAEDLPYQSSLASWNYNTNITEENVQNNAGDGKWSAPLKEQST	60
Db	18	STTIEQAKTFLDKFNHEAEDLPYQSSLASWNYNTNITEENVQNNAGDGKWSAPLKEQST	77
Qy	61	LAQWYPLQIBQNLTVKQLQALQOQSSVLSEDKSKRLNTILNTMSTIYSTGKVCNPDNP	120
Db	78	LAQWYPLQIBQNLTVKQLQALQOQSSVLSEDKSKRLNTILNTMSTIYSTGKVCNPDNP	137
Qy	121	QECLLLEPGLNEIWMANSLDYNERLWAWESWRSEVGKQLRPLYEYVVLKNEMARANHYED	180
Db	138	QECLLLEPGLNEIWMANSLDYNERLWAWESWRSEVGKQLRPLYEYVVLKNEMARANHYED	197
Qy	181	YGDYWRGDEVNGVDGYDSRGQLIBDVEHTFBEIKPLYEHLHAYVRAKLMNAYPSYISP	240
Db	198	YGDYWRGDEVNGVDGYDSRGQLIBDVEHTFBEIKPLYEHLHAYVRAKLMNAYPSYISP	257
Qy	241	IGCLPAHLIGDMKGRFWNTNLSLTVPGQKPNIDVTDAMVDQAWDAQRIKPEAEKFFVSU	300
Db	258	IGCLPAHLIGDMKGRFWNTNLSLTVPGQKPNIDVTDAMVDQAWDAQRIKPEAEKFFVSU	317
Qy	301	GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDDFLTAHHEMGH	360
Db	318	GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVMTDDDFLTAHHEMGH	377
Qy	361	IQYDMAYAAQPFLLRNGANEGPHEAVGEIIMSLSAATPKHLKSIGLLSPDFOEDNETEINF	420
Db	378	IQYDMAYAAQPFLLRNGANEGPHEAVGEIIMSLSAATPKHLKSIGLLSPDFOEDNETEINF	437
Qy	421	LLKQALTVIGTLLPPTMYMLEKRWMMVFKEIIPKQOAWKKWHEMKREIIVGVVSEPUPHDETIC	480

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Db 438 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGIPKQDQMMKKWEMKREIVGVVPEVPHDETYC 497
QY 481 DPASLPHVNSDYSFIRYRTLYQFOFQALCOAAKHEGPLHKCDISNSTEAGQKLENNL 540
Db 498 DPASLPHVNSDYSFIRYRTLYQFOFQALCOAAKHEGPLHKCDISNSTEAGQKLENNL 557
QY 541 RLKGSFPWTALLENVVGAKMNVRLPLNLYFEPLFTWLKQDNKNSFVGNSTWSPY 595
Db 558 RLKGSFPWTALLENVVGAKMNVRLPLNLYFEPLFTWLKQDNKNSFVGNSTWSPY 612

RESULT 2
S05238
peptidyl-di-peptidase A (EC 3.4.15.1) precursor, testicular splice form - human
N:Alternate names: angiotensin I-converting enzyme (ACE); CD143; dipeptidyl carboxypeptid
C:Species: Homo sapiens (man)
C>Date: 30-Sep-1991 #sequence_revision 02-Jul-1998 #text_change 09-Jul-2004
R:Rattion, A.L.; Soubrier, F.; Allegri, J.; Hubert, C.; Corvol, P.; Alhenc-Gelae, F.
FEBS Lett. 252, 99-104, 1989
A:Title: The testicular transcript of the angiotensin I-converting enzyme encodes for th
A:Reference number: S05238; MUID:89338720; PMID:2547653
A:Accession: S05238
A:Molecule type: mRNA
A:Residues: 1-732 <LAT>
A:Cross-references: UNIPROT:P22966; UNIPARC:UPI000002DB19; EMBL:X16295; NID:G28264; PIDN
R:Ehlers, M.R.W.; Fox, E.A.; Strzydom, D.J.; Riordan, J.F.
Proc. Natl. Acad. Sci. U.S.A. 86, 7741-7745, 1989
A:Title: Molecular cloning of human testicular angiotensin-converting enzyme: the testie
A:Reference number: A33979; MUID:90046671; PMID:2554286
A:Accession: A33979
A:Molecule type: mRNA
A:Residues: 1-732 <EHL>
A:Cross-references: UNIPARC:UPI000002DB19; GB:M26657; NID:G338666; PIDN:AAAG0611.1; PID:
A:Experimental source: clones R1.2 and T88
A>Note: neither the complete nucleic acid sequence nor the complete translation are show
C:Comment: For the renal and pulmonary splice form, see PIR:A31759.
C:Genetics:
A:Gene: GDB:DCPI; ACE
A:Cross-references: GDB:119840; OMIM:106180
A:Map position: 17q23-17q23
C:Function:
A:Description: catalyzes the hydrolysis of dipeptides from the carboxyl end of polypepti
C:Superfamily: mammalian peptidyl-di-peptidase A
A:Keywords: alternative splicing; glycoprotein; metalloproteinase; peptidyl-dipeptide hyd
F:1-21/Domain: signal sequence #status predicted <SIG>
F:22-732/Product: peptidyl dipeptidase I #status predicted <MAT>
F:686-702/Domain: transmembrane #status predicted <TRM>
F:103,121,140,186,368,617,651/Binding site: carbohydrate (Aan) (covalent) #status predic
F:414,418,434/Binding site: zinc, catalytic (His, His, Glu) #status predicted
F:415/Active site: Glu #status predicted

Query Match 41.3%; Score 1335; DB 1; Length 732;
Best Local Similarity 41.9%; Pred. No. 6.6e-88;
Matches 254; Conservative 116; Mismatches 202; Indels 34; Gaps 9;

QY 2 TTEEQAKTFLDKFNHEADLFFQSSLASNNYNTITEE-----NVQNNNAGDKWSA 53
Db 70 TDEARASKEVEYDRTSQVNNYAEANNNYNTINTTETSKILLQNNQIANHT----- 123

QY 54 FLKEQSTLAQMPLOBIQNLTKLQALQALQNGSSVLSDSKSKRLNTIINTSTIYSTGK 113
Db 124 --LKYGTOARKFDVNLQNTTKRIKKVQDLERAAALPAQELEEYKILLDMETIYSAT 181

QY 114 VCNPNPOBCLLEPGELNEMANSIDYNERLWANSRSEVGKQLRPLYEYVVLKNEWA 173
Db 182 VCHPNG--SCLOLEPLTNWATSRKYEDLLWAEGRDKAGRAILQPPKVELINQAA 239

QY 174 RANHYEDYGVNRGVNGVDGYDSRQLLEDVEHTEFKPLYEHLHVAVRKLMA 233
Db 240 RLVNGYVDAGDSWRSMYETPSLE-----QDLERFQELQPLYLNLHVAVRALHRH 289

QY 234 Y-PSYISPIGCLPAHLGLDMGRFWTNLYSLTVFPGQKNIDVTDAMVDQAWDAORIFKE 292
```

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Db 290 YGAQHINLEGPIPAHLGNMMAQTWSNIYDLVVPPSPASMDTTEAMLKQGWTPRRMFKE 349
QY 293 AEKPFVSVGLPNMTQGFWENSHLTDPGNVQKAVCHPTANDLGKG--DFRILMCTKTYMDDF 351
Db 350 ADDFTSLGLLFPVPEFWKSMLEKPTDGRVCHASAWDFYNGKDFRIKQCTTVNLEDL 409
QY 352 LTAHHEMGHIQVDMAAAQPFLLRNGANEHFEAETMSLSAAATPKHLKSTGLLSPDQF 411
Db 410 VVAHHEMGHIQYFMQYKDLPVALREGANPGFHEALGDVIALSVSTPKHLHSLNLLSSGG 469
QY 412 EDNETEINFLKQALTIIVGTLPTFTYMLEKRWMMVFKGIPKQDQMMKKWEMKREIVGVV 471
Db 470 SD-EHDINFLMKALDKIAPFSLVDQWRVRVDSGITTKENYNQEWLSRLKYQGLCP 528
QY 472 PVPHDETYCDASLPHVNSDYSFIRYRTLYQFOFQALCOAAKHEGPLHKCDISNSTE 531
Db 529 PVPRTQSGDFDCAKHPFSSVPYIRYFVSFIQFQHEALCOAAAGHTGPLHKCDIYQSK 588
QY 532 AGQKLFNMLRLKGSFPWTALLENVVGAKMNVRLPLNLYFEPLFTWLKQDNK--NSFVGW- 588
Db 589 AGQRLATAMKLGSRPPEAMQLITQDPNMSASAMLSYFKPLLDWLRTENELHGEKLGWP 648
QY 589 STDWSP 594
Db 649 QYNWTP 654

RESULT 3
A31759
peptidyl-di-peptidase A (EC 3.4.15.1) precursor, renal and pulmonary splice form - human
N:Alternate names: angiotensin I-converting enzyme (ACE); CD143; dipeptidyl carboxypepti
C:Species: Homo sapiens (man)
C>Date: 07-Jun-1990 #sequence_revision 02-Jul-1998 #text_change 09-Jul-2004
C:Accession: A31759; PQ0004
R:Soubrier, F.; Alhenc-Gelae, F.; Hubert, C.; Allegri, J.; John, M.; Tregear, G.; Corv
Proc. Natl. Acad. Sci. U.S.A. 85, 9386-9390, 1988
A:Title: Two putative active centers in human angiotensin I-converting enzyme revealed by
A:Reference number: A31759; MUID:89071703; PMID:2849100
A:Accession: A31759
A:Molecule type: mRNA
A:Residues: 1-1306 <SOU>
A:Cross-references: UNIPROT:P12821; UNIPARC:UPI000002B9AD; GB:J04144; NID:G178285; PIDN:
A:Experimental source: kidney
A>Note: parts of this sequence, including the amino end of the mature protein, were dete
R:Takeuchi, K.; Shimizu, T.; Ohishi, N.; Seyama, Y.; Takaku, F.; Yotsumoto, H.
J. Biochem. 106, 442-445, 1989
A:Title: Purification of human lung angiotensin-converting enzyme by high-performance li
A:Reference number: PQ0004; MUID:90110025; PMID:2558109
A:Accession: PQ0004
A:Molecule type: protein
A:Residues: 'XX', 32-34, 'G', 36-37, 'X', 39-41, 'R', 43-46 <TAK>
A:Cross-references: UNIPARC:UPI00000172A3D
A:Experimental source: lung
A:Comment: This splice form is found in many tissues, in particular kidney and lung vascu
C:Genetics:
A:Gene: GDB:DCPI; ACE
A:Cross-references: GDB:119840; OMIM:106180
A:Map position: 17q23-17q23
C:Function:
A:Description: catalyzes the hydrolysis of dipeptides from the carboxyl end of polypeptic
A>Note: plays a role in the control of blood pressure by catalyzing the conversion of ang
C:Superfamily: mammalian peptidyl-di-peptidase A
C:Keywords: alternative splicing; blood pressure control; glycoprotein; kidney; lung; met
F:1-29/Domain: signal sequence #status predicted <SIG>
F:30-1306/Product: peptidyl dipeptidase I #status predicted <MAT>
F:1260-1276/Domain: transmembrane #status predicted <TRM>
F:38,54,74,111,146,160,318,445,509,523,677,695,714,760,942,1191,1225/Binding site: carbo
F:390,394/Binding site: zinc (His) #status predicted
F:988,992,1008/Binding site: zinc, catalytic (His, His, Glu) #status predicted
F:989/Active site: Glu #status predicted

Query Match 41.3%; Score 1335; DB 1; Length 1306;
```


C:Keywords: alternative splicing; blood pressure control; membrane protein; peptidyl dipeptidase I
F:1-34/Domain: signal sequence #status predicted <SIG>
F:35-1312/Product: peptidyl dipeptidase I #status predicted <MAT>

Query Match 41.3%; Score 1334; DB 1; Length 1312;
Best Local Similarity 42.6%; Pred. No. 1.9e-87;
Matches 255; Conservative 112; Mismatches 213; Indels 18; Gaps 7;

2 TIEBQAKTFIDKFNHEADLPYQSSLASNNYNTNITTEENVQNMNAGDKWSAFLEQSTL 61
649 TDEAKADRFVEEYDRTQAVLLNEVAENWQNTNITIEGSKILLEKSTEVSNHTLKYGTR 708
62 AQMYPLQEIQNLTVKLOLQALQQNGSSVLGSDKSKRLNTILNTWMTTISTYTKVCNPNPQ 121
709 AKTFDVSFNQSSIKRIKKLQNLDRVLPPEKEEYNQILLDMETYSLSNICYTG-- 766
122 ECLILEPLNEIMANSIDYNERLWAWESWSEVKQRLRPLYEEYVVLKNEMARANHYEDY 181
767 TCMLEPLDNTMNAATSKRYEELLWAWKSRDKGRALPFPFKYVEFSNIAKILNGYTD 826
182 GDYWRGDYVNGVDGYDSRGQLTDEHTEPEETKPLYEHLHAYVRAKLMAYPS-YISP 240
827 GDSWRSYVESDNLE-----QDLEKLYQELQPLYLNLHAYVRSLSRHYGSEVNL 876
241 IGCPLPAHLGLDMGRFNTNLSYLVTPFGQKPNIDVTDAVDVQAWDAQRIKFAEKFFVS 300
877 DGPPIPAHLGLNMAQTWSNIYDLVAPFSPAPNIDATEAMIKQGWTPRIRFKEADNFTSL 936
301 GLPNWTOGFVENSMLTDPGNVQKAVCHPTAWDLKGG-DEFILMCTKYTMDDLTAHHEMG 359
937 GLLPVPPEFVFNKSMLEKPTDGRVVVVCHPSAWDFNGKDFRIKQCTSNMBSDLVIAHHEMG 996
360 HIQYDMAYAAQFFLLRRNGANEGFHEAVGEIMSLSAATPKHLKSLGLSPQFQEDNTEIN 419
997 HIQYFMQYKDLPVTFREGANPGFHEAIGDMALSVSTPKHLYSLNLLSTB-GSGYEYDIN 1055
420 FLLKQALITVGTLPFTMLEKWRMVFKEIPIKQWKKWEMKREIVGVVPEVPHDETY 479
1056 FLKQVALDKIAFIPESFLIDQWRVRVFGSITKENYQWMSLTKYQGLCPVPRSQGD 1115
480 CDPASLPHVSNDSYFIRYRTTLVQFOFQEAALCOAAKHGEPHLHKCDISNTEAGOKLFNM 539
1116 FDPQSKFHPANVPVYRVYFVFIQFQFHEALCRAAGHTGPLHKCDIYQSKKAGKLADA 1175
540 LRLGKSPFTWLTALENVVGAKNMVRPLNLYPEPLFTWLKDNK--NSFVGH-STDNSP 594
1176 MKLGYSKPWPEAMKLIITGQPNWSASAMNYPFKPLTEWLVTEENRRHGHTGLGWPEYNWAP 1233

RESULT 6
JC2038
peptidyl-di-peptidase A (EC 3.4.15.1) - rat
N;Alternate names: angiotensin converting enzyme; Kininase II
C;Species: Rattus norvegicus (Norway rat)
C;Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 09-Jul-2004
C;Accession: JC2038
R;Koike, G.; Krieger, J.B.; Jacob, H.J.; Mukoyama, M.; Pratt, R.E.; Dzau, V.J.
Biochem. Biophys. Res. Commun. 198, 380-386, 1994
A;Title: Angiotensin converting enzyme and genetic hypertension: Cloning of rat cDNAs and characterization of the complementary DNA
A;Reference number: JC2038; MUID:94121658; PMID:8292044
A;Accession: JC2038
A;Molecule type: mRNA
A;Residues: 1-1313 <NOI>
A;Cross-references: UNIPROT:P47820; UNIPARC:UPI000016928F; GB:U03734; NID:g437289; PIDN:
A;Note: The authors translated the codon ACC for residue 159 as Tyr
C;Comment: This enzyme is a zinc-containing carboxy peptidase that cleaves angiotensin I
C;Superfamily: This enzyme plays a critical role in blood pressure homeostasis and is the catalytic subunit of the angiotensin converting enzyme complex
C;Comment: This enzyme plays a critical role in blood pressure homeostasis and is the catalytic subunit of the angiotensin converting enzyme complex
C;Keywords: alternative splicing; peptidyl dipeptide hydrolase; transmembrane protein; z
F;393-400/Region: catalytic #status predicted
F;1264-1284/Domain: transmembrane #status predicted <TM>

Query Match 40.5%; Score 1310; DB 1; Length 1313;

Best Local Similarity 42.0%; Pred. No. 1e-85;
Matches 251; Conservative 115; Mismatches 214; Indels 18; Gaps 7;

QY 2 TIERQAATFLDKNHEADLFYQSGLASWNTYNITTEENVQNMMNAGDKWSAFLKEQSTL 61
Db :|::||:::||::||::||::||::||::||::||::||::||::||:
650 TDEAKANRFVEEYDRTAKVLANVEAAWHNYNTNIIEGSKILLKCNKEVSNNHTLKYGTV 709
QY 62 AQMYPLQEIQNLTKVLQLAQLOQGSSVLSDKSRLRINTILMTSTIYSTGVKVCNDPNPQ 121
Db :|::||:::||::||::||::||::||::||::||::||::||:
710 AKTDFVNFSNQSTIKRIKKVQNVDRALVPNELEYNOILLDMETTYSVANVCYTNG-- 767

QY 122 ECLILEFGLINEIMSLDYNERLNAMESWRSEVGKQLRPPLYEEYVVKEMEBARANEHVEDY 181
Db :|::||:::||::||::||::||::||::||::||::||::||:
768 TCLESLEPDLTINMATSRYKEYSELLMWKSWRDKVGRAILPFPPFKYVDVFNKIACLKGYSDA 827

QY 182 GDYWVRGDYEVNGVDGYDSRGOLIEDVBETHPEIRKPLYEHHLHYAVRAKLMLNAYPS-YISP 240
Db :|::||:::||::||::||::||::||::||::||::||::||:
828 GDSWRSYESDDL-----QDLKELVQLLPYLINLHAYVRSLSRHVGSYYNL 877

QY 241 IGCPLAHLLGDMGRFWTNLYSLTPFGQPKNIDVTMDAMVDQAWDAQORIFKEAEKFVSU 300
Db :|::||:::||::||::||::||::||::||::||::||::||:
878 DGPPIPHALLGNMAQTNSNIVLDVAFFSPAPSIDATEAMI KQGWTPRRIFKEADNFETS L 937

QY 301 GLPNNTOGFWENSMILTDPGNVQKAVCHPTAMDLGKG-DPRILMCITYKTWDDEFATAHHMG 359
Db :|::||:::||::||::||::||::||::||::||::||::||:
938 GLFPVPEFWNKMLEKFTDGREVVCHASANDFYNGKDFRIKQCTSVNMHEELVIANHENG 997

QY 360 HIQYDMAAYAAPFLLRNCANGSGFEHAVGEIMSLSAATPKHKLSIGLLSDFPOEDNETEIN 419
Db :|::||:::||::||::||::||::||::||::||::||::||:
998 HIQTFMQYKDLLPVTFREGANGFHGAIGDVALSSTPKHLHSLLNLSSE-GSGYEHDIN 1056

QY 420 FLLKOALTITGTLPFTVMLEKWRMVFGEIPKDQQMKWKWMKBEIVGVVEVPVPHDET Y 479
Db :|::||:::||::||::||::||::||::||::||::||::||:
1057 FLMKVALDKIAFI PFSYLIIDQMRWFVDSITKENYNGEWMSLRLLKY QGLCPVPPRS QGD 1116

QY 480 CDPASLFHVSNDSYFIRYYTTTLTQFOQBALCOAAKHGHPHKCDISNSTEQGXKLFWM 539
Db :|::||:::||::||::||::||::||::||::||::||::||:
1117 FDPGSKHFVPANVPYIRFYIFIIOQOPHEALCRAAGHTGPLYCDDIYO SKERAGKLADA 1176

QY 540 LRLLCKSEPWTLALENVVGAKMNVRPLANYPEPLTWLKDQNK--NSFVGM-STDWSP 594
Db :|::||:::||::||::||::||::||::||::||::||::||:
1177 MKUGYSKOMPEAMKIITGOPNNASAINYPKPLEWLVTENRRHGTEL GWPEYTWTP 1234

RESULT 7
JC2489
peptidyl-dipeptidase A (EC 3.4.15.1) - chicken
N/Alternate names: angiotensin converting enzyme
C/Species: Gallus gallus (chicken)
C/Date: 16-Mar-1995 #sequence_revision 26-May-1995 #text_change 09-Jul-2004
C/Accession: JC2489
R/Esther, C.R.; Thomas Jr., K.E.; Bernstein, K.E.
Biochem. Biophys. Res. Commun. 205, 1916-1921, 1994
A>Title: Chicken lacks the testis specific isozyme of angiotensin converting enzyme found
A/Reference number: JC2489; PMID:95110342; PMID:7811282
A/Accession: JC2489
A/Molecule type: mRNA
A/Residues: 1-1193 <SUP>
A/Cross-references: UNIPROT.Q10751; UNIPARC.UPI000012521B; GB:I40175; NID:g685168; PIDN:/
C/Comment: This enzyme is a zinc dependant dipeptidyl carboxypeptidase that cleaves a vari
C/Superfamily: mammalian peptidyl-di-peptidease A
C/Keywords: metal binding; peptidyldipeptide hydrolase; zinc
F/316,331,914,929/Binding site: zinc, catalytic (Glu, His, Glu, His) #status predicted

Query Match 40.5% Score 1307; DB 2; Length 1193;
Best Local Similarity 42.3%; Pred. No. 1.4e-85;
Matches 254; Conservative 106; Mismatches 216; Indels 24; Gaps 8;

QY 4 BEQAQTFLLDENHEADLFYQSGLASWNTYNITTEENVQNMMNAGDKWSAFLKEQSTLAQ 63
Db :|::||:::||::||::||::||::||::||::||::||::||:
544 EAQAKEFUSESTADEVVNNAYTEASWEISNTINITHNKCELMLEKANLSKHTLEYGNRAR 603

QY 64 MYPLQEIQNLTKVLQLAQLOQGSSVLSDKSRLRINTILMTSTIYSTGVKVCNDPN---P 120

Db	604	QFPPSFDQETVTRILNKLSVLERAAIPDELDKEYNTLLSDMETTVSVAKVCRENNTFHP	663
Qy	121	QECLEPGLNEIMANSLDYNRLAWESWRSEVGKQLRPLRYEYVVLKXNEMARANHYED	180
Db	664	-----LDPDLTDILATSRDYNELLPAWKGDWASGAKIKDYKRYVELSKAAVLNGYTD	718
Qy	181	YGDYWRGDEYVNGVDYDYSRGQLIDVHTPEEIKPLIEHLHAYVRAKLMAY-PSYIS	239
Db	719	NGAYWRSLEYETTFE-----BDLERLYLQQLPLYLNLHAYVRRALYNKYGAHIS	768
Qy	240	PIGCLPAHLGDMGWFNTNLSLTVPGQKNIDVTDAMVDQDAQRIKFAEKFFVS	299
Db	769	LKGPIPAHLGNNWASGNSIFDLVNPFPDATKYDPAWKQGGWTPKWMFESDRFFTS	828
Qy	300	VGLPNNTOGFWNSMLTDPGNVQKAVCHPTAMD-CKGDFRILMCKTVKTMDDFLTAHHEM	358
Db	829	LGLIPMPQFQWDSKMEKPADGREVVCHASAWDFYNRKDFRIKQCTVVMDDLIIVHEM	888
Qy	359	GHIQYDMAAQPFLRNGANEGFHAEGEIMSLSAATPKHLKSGILLSPDFQEDNETEI	418
Db	889	GHVQVPLQTMQDPISRFDGANPGFHAIGDVNALSSTPKHLHSINLLD-QVTENBESDI	947
Qy	419	NFLKKAALITVGLPTMYLKWVNVFKEGPEIKQDMKKWEMKEIIVGVVPEVPHDET	478
Db	948	NYLMSIALDKIAFLPGYGLMDQWRKVFQGRKEDESYNQWNNRLUKYQGLCPPVPSRD	1007
Qy	479	YCDPASLFHVSNDYSIRYTRTYQFQEQALCOAKHEGPHLKCDISNSTEAGOKLPN	538
Db	1008	DFDPGAKFHI PANVPYIRYFVSFVIOFQFQALCKAAGTGPHLTCDIYQSKAAGKLGCD	1067
Qy	539	MURLKSEPTWTLAENVGAKNNVRPLNRYPEPLFTWL--KDQNKNSFVGM-STDWSPY	595
Db	1068	AMKLGSKWPEAMQIITGQPNMSAELMSYEPLMTLVKXNTENGELVGPYSWTPE	1127
RESULT 8			
A34402			
peptidyl-dipeptidase A (EC 3.4.15.1) precursor, testicular - rabbit			
N:Alternate names: angiotensin I-converting enzyme; dipeptidyl carboxypeptidase I; pepti			
C:Species: Oryctolagus cuniculus (domestic rabbit)			
C:Date: 10-Sep-1999 #sequence revision 10-Sep-1999 #text_change 09-Jul-2004			
C:Accession: A34402; A60724; A36232; C18700			
R:Kumar, R.S.; Kusari, J.; Roy, S.N.; Soffer, R.L.; Sen, G.C.			
J. Biol. Chem. 264, 16754-16758, 1989			
A:Title: Structure of testicular angiotensin-converting enzyme. A segmental mosaic isozy			
A:Reference number: A34402; MUID: 89380303; PMID: 2550457			
A:Accession: A34402			
A:Molecule type: mRNA			
A:Residues: 1-737 <KUM>			
A:Cross-references: UNIPROT:P22968; UNIPARC:UPI000004A558; GB:J05041; NID:g164744; PIDN:			
R:Sen, G.C.; Thekkumkara, T.J.; Kumar, R.S.			
J. Cardiovasc. Pharmacol. 16(Suppl. 4), S14-S18, 1990			
A:Title: Angiotensin-converting enzyme: structural relationship of the testicular and the			
A:Reference number: A60724; MUID: 91155372; PMID: 1705622			
A:Accession: A60724			
A:Status: translation not shown			
A:Molecule type: mRNA			
A:Residues: 73-173 <SEN>			
A:Cross-references: UNIPARC:UPI0000172A3P			
A:Note: Identical sequences were obtained for mRNAs from lung and testes			
R:Chen, Y.N.P.; Riordan, J.F.			
Biochemistry 29, 10493-10498, 1990			
A:Title: Identification of essential tyrosine and lysine residues in angiotensin convert			
A:Reference number: A36232; MUID: 91104959; PMID: 2176870			
A:Accession: A36232			
A:Molecule type: protein			
A:Residues: 154-160/236-242 <CHE>			
A:Cross-references: UNIPARC:UPI0000172A40; UNIPARC:UPI0000172A41			
R:Twata, K.; Lai, C.Y.; El-Dorry, H.A.; Soffer, R.L.			
Biochem. Biophys. Res. Commun. 107, 1097-1103, 1982			
A:Title: The NH2- and COOH-terminal sequences of the angiotensin-converting enzyme isozym			
A:Reference number: A90107; MUID: 83048249; PMID: 62991514			

A:Accession: C18700			
A:Molecule type: protein			
A:Residues: 33-35, 'SN', 38-39, 'SS', 'PAEL', 737 <IWA>			
A:Cross-references: UNIPARC:UPI0000172A42; UNIPARC:UPI0000172A43			
A:Note: several of the amino acids in reported are tentative			
C:Comment: The pulmonary and testicular isoforms of this enzyme differ substantially in l			
gests that the two isoforms arise by alternative splicing of one gene.			
C:Superfamily: mammalian peptidyl-dipeptidase A			
C:Keywords: alternative splicing; peptidyl-dipeptide hydrolase; testis; transmembrane prot			
Query Match			
Best Local Similarity 41.0%; Score 1281; DB 1; Length 737;			
Matches 249; Conservative 110; Mismatches 211; Indels 38; Gaps 10;			
Qy	2	TIEQAKTFLDKFNHEADLFYQSSLASNNYNTNITE-----NVQNNNN--AGDKW	51
Db	75	TDEASRSFVEYDRSFQAVNVEYAEANNVNTNITTEASKILLQKNMQIANHLLTYGNW	134
Qy	52	SAFLKEQSTLAQMYPLQEIQNLTVKLQALQOQSSVLSSEKSKRLANTILNTMTSTYST	111
Db	135	-----ARRFDVSNFQNAISRRIKKVQODLQRAVLPVKELBEYQIILDMETIYSV	184
Qy	112	GRVCNPDNPQECLELLEPGLNEIMANSLDYNRLAWESWRSEVGKQLRPLRYEYVVLKNE	171
Db	195	ANVCERVDG--SCLQLEPDLTNLMATSRKYDELLWVWTSWRDKVGRAILPYFPKYVEFTNK	242
Qy	172	MARANHYSDYGYWRGDEYVNGVDYDYSRGQLIDVHTPEEIKPLIEHLHAYVRAKLM	231
Db	243	AARLNGYVDAGDSWRSMTETPTLE-----QDLERLFOELQPLVNLHAYVGRALH	292
Qy	232	NAY-PSYISPTGCLPAHLGDMGWFNTNLSLTVPGQKNIDVTDAMVDQDAQRIK	290
Db	293	RHYGAOHINLEGPIPAHLGNNWAGTWSNIYDLVAFPSASTMDATEAMIKOGWTPRRMF	352
Qy	291	KEAEKFFSVGLPNNTOGFWNSMLTDPGNVQKAVCHPTAMDGLKG-DPRILMCKTVMD	349
Db	353	BEADKFFISLGLLPVPPEFWNKSMLKEPTDGREVVCHASAWDFYNGKOPRIKQCTVANE	412
Qy	350	DEFLTAHEMGHIQYDMAAQPFLRNGANEGFHAEGEIMSLSAATPKHLKSGILLSPD	409
Db	413	DLVVVHEMGHIQYPMQYKDLPVALLREGANPGFHAIGDVLALSSTPKHLHSINLLSE	472
Qy	410	FOEDNETINELLKQALITVGLPTMYLKWVNVFKEGPEIKQDMKKWEMKEIIVGV	469
Db	473	-GGGYEHDFINFLKQALDKIAFIPPSYLVDEWRVFDGSIKENYNGEWSRLUKYQGL	531
Qy	470	VEPVPHDETCDPASLFHVSNDYSIRYTRTYQFQEQALCOAKHEGPHLKCDISNS	529
Db	532	CPPAPRSQDDPFGAKFHIPSSVPYIRYFVSFVIOFQFQALCKAAGTGPHLTCDIQS	591
Qy	530	TEAGOKLFNMLRLKSEPTWTLAENVGAKNNVRPLNRYPEPLFTWLKQDN--KNSFVG	587
Db	592	KEAGKRLADAMKLGYSKWPPEAMKVITGQPNMSASAMWNYPKPLMDWLLTENGHRGEKLG	651
Qy	588	W-STDWSP	594
Db	652	WPQYTWTP	659
RESULT 9			
S35484			
peptidyl-dipeptidase A (EC 3.4.15.1) precursor, pulmonary splice form - rabbit			
N:Alternate names: angiotensin-converting enzyme; dipeptidyl carboxypeptidase I; kininase			
C:Species: Oryctolagus cuniculus (domestic rabbit)			
C:Date: 10-Sep-1999 #sequence revision 10-Sep-1999 #text_change 31-Dec-2004			
C:Accession: S35484; A23455; A18700; A38655; A49726; S17509			
R:Thekkumkara, T.J.; Livingston III, W.; Kumar, R.S.; Sen, G.C.			
Nucleic Acids Res. 20, 683-687, 1992			
A:Title: Use of alternative polyadenylation sites for tissue-specific transcription of t			
A:Reference number: S35484; MUID: 92178960; PMID: 1311831			
A:Accession: S35484			
A:Molecule type: mRNA			
A:Residues: 1-1309 <THE>			

A;Cross-references: UNIPROT:P12822; UNIPROT:Q9TRW7; UNIPARC:UPI00000172A44; EMBL:X62551
R;Iwata, K.; Blacher, R.; Soffer, R.L.; Lai, C.Y.
Arch. Biochem. Biophys. 227, 188-201, 1983
A;Reference number: A23455; MUID:84051289; PMID:6314908
A;Accession: A23455
A;Molecule type: protein
A;Residues: 34-47, 'N', 49-55 <IWA>
A;Cross-references: UNIPARC:UPI00000172A45
A;Experimental source: lung
R;Iwata, K.; Lai, C.Y.; El-Dorry, H.A.; Soffer, R.L.
Biochem. Biophys. Res. Commun. 107, 1097-1103, 1982
A;Title: The NH2- and COOH-terminal sequences of the angiotensin-converting enzyme isozym
A;Reference number: A90107; MUID:83048249; PMID:6291514
A;Accession: A18700
A;Molecule type: protein
A;Residues: 34-44;754-755, 'L', 757 <IW2>
A;Cross-references: UNIPARC:UPI00000172A46; UNIPARC:UPI00000172A47
R;Kumar, R.S.; Thekkumkara, T.J.; Sen, G.C.
J. Biol. Chem. 266, 3854-3862, 1991
A;Title: The mRNAs encoding the two angiotensin-converting isozymes are transcribed from
A;Reference number: A38655; MUID:91139683; PMID:1847388
A;Accession: A38655
A;Molecule type: DNA
A;Residues: 1-88 <KUM>
A;Cross-references: UNIPARC:UPI00000172A48; GB:M58579
R;Ramchandran, R.; Sen, G.C.; Misono, K.; Sen, I.
J. Biol. Chem. 269, 2125-2130, 1994
A;Title: Regulated cleavage-secretion of the membrane-bound angiotensin-converting enzym
A;Reference number: A49726; MUID:94124568; PMID:8294466
A;Accession: A49726
A;Status: preliminary
A;Molecule type: protein
A;Residues: 1236-1258 <RAM>
A;Cross-references: UNIPARC:UPI00000172A49
A;Experimental source: testis
R;Kirley, J.L.
Biochem. J. 278, 375-380, 1991
A;Title: The Mg(2+)-ATPase of rabbit skeletal-muscle transverse tubule is a highly glyco
A;Reference number: S17509; MUID:91378880; PMID:1654980
A;Accession: S17509
A;Status: preliminary
A;Molecule type: protein
A;Residues: 34-55 <KIR>
A;Cross-references: UNIPARC:UPI00000870D8
C;Comment: This enzyme converts angiotensin I to angiotensin II in presence of divalent
ver, the enzyme has been found also in renal tubules and intestinal mucosa.
C;Keywords: alternative splicing; blood pressure control; chloride; glycoprotein; intest
F;1-33/Domain: signal sequence #status predicted <SIG>
F;34-11309/Product: peptidyl-diesterase A, pulmonary #status experimental <MAT>
P;59,79,150,322,448,512,680,698,717,945,1194/Binding site: carbohydrate (Asn) (covalent)

Query Match 39.68; Score 1281; DB 1; Length 1309;
Best Local Similarity 41.08; Pred. No. 1.2e-83;
Matches 249; Conservative 110; Mismatches 211; Indels 38; Gaps 10;

Qy 2 TIEQAKTFLDKFNHEAEDLFYOSSLASWNYNTNITEE-----NVQNMNN--AGDKW 51
Db 647 TDEASRFFVEYDRSFQAVNVEAYEANNWYNTNITEASKILLQNMQIANHTLYGNW 706
Qy 52 SAFLKEOSTLAQMPYLPQEIQNLTQVLQALQNGSSVLSEDSKKRLNTILNTMTSYST 111
Db 707 -----ARRFDVSFNQATSKRIKKVQDLQRAVLPAVKELEEVNQILLDMETIYSV 756
Qy 112 GKVCNPDNPQECILLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLIYEVVYVKN 171
Db 757 ANVCRVDG--SCQLQEPDLTNLMATSKRYDELLWMTWSMRDKVGRAILPFPKPYVEFTNK 814
Qy 172 MARANHYEDYGDYWRGVEYNGVDGYSRGQLIEDVEHTFERIKPLYEHLHAYVRAKLM 231
Db 815 AARLNGYVDAGDSWRSWYETPTLE-----QDLERLFQELQPLYNLHAYVGRALH 864
Qy 232 NAY-PSYISPIGCLPAHLLGDMMGRFWNTLYSLVTPFGQKPNIDVTDAVQDAQRIF 290

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Db 198 TDGAEVWLDEYE-----DATFEOLEAFEDIKPLVDQVHGYYRRLNKFYGD 246
Qy 237 YISPIGCLPAHLIGDMWGRFTWNLVSLTVPFGQKNIDVTDAMVDQAWDAQRIKFAEAKF 296
Db 247 VVSKTGCLPMLHLLGNMAQWMSIADIVSPFPEKPLVDVSDENVAQGYTPLKMFQMGDDF 306
Qy 297 FVSVGLPNTQGFWNSMLTDPGNQKAVCHPTAWDLG-KGDPRIILMCTKVTWDDFLTAH 355
Db 307 POSMGLKGLPQEFWDSKILEKPDGDRDLVCHASAWDFYLTDDVRIKQCRTVQDQFFTVH 366
Qy 356 HEMGHLYQDMAYAAQPFLLRNGANEGFHEAVGIMSLSAATPKHLKSGILLSPDFQEDNE 415
Db 367 HEMGHLYQFLOYQHFPVIRGTGANGPGFHEAVGVDLSVSTPKHLRVLGK-NYVSDNE 425
Qy 416 TEINFLKQALITVGLTPTMYLWKRMVVFKEIPKQDMKMKWEMKREIIVGVVEPVPH 475
Db 426 ARINQLFLTALDKIVLPFAFTMDKYRWALFRQADKSEWNCFAFWKLREYSIGIRPPVR 485
Qy 476 DETYCDPASLPHVNSDYSIRYTRTYQFOQFQALCOAA-----KHGPHLHKCDISN 529
Db 486 TEKDFAPAKYHVSADVEYLRVLSFIIQFPYKACITAGEVVNPQTEYPLDNCDIYS 545
Qy 530 TEAGOKLFWMLRGKSEPTLALENVVGAKNNVRPLLNYERPLFTWLK 578
Db 546 KEAGKLFENWLSLGAKPMDALAEAFNGERTMTGKAIAEYFBLRWLE 594

RESULT 11
JC5374
angiotensin-converting enzyme-related protein - fruit fly (Drosophila melanogaster)
C:Species: Drosophila melanogaster
C:Date: 28-May-1997 #sequence_revision 18-Jul-1997 #text_change 09-Jul-2004
C:Accession: JC5374
R:Taylor, C.A.M.; Coates, D.; Shirrae, A.D.
Gene 181, 191-197, 1996
A:Title: The Acer gene of Drosophila codes for an angiotensin-converting enzyme homolog
A:Reference number: JC5374; MUID:97128790; PMID:8973330
A:Accession: JC5374
A:Molecule type: mRNA
A:Residues: 1-630 <P>A>
A:Cross-references: UNIPROT:Q24222; UNIPARC:UPI0000075442; EMBL:X96913; NID:g1405881; PI
C:Genetics:
A:Gene: Acer
C:Superfamily: mammalian peptidyl-dipeptidase A

Query Match 31.8%; Score 1027.5; DB 2; Length 630;
Best Local Similarity 36.0%; Pred. No. 6.7e-66;
Matches 215; Conservative 110; Mismatches 250; Indels 23; Gaps 9;

Qy 6 QAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNNMAGDKWSAFLEKQSTLAQMY 65
Db 33 EARRPELENEQLRRRFHEFLSGYNYNTNTEANRQAMIEVYARNALNKLKLAQKISS 92
Qy 66 PLOEIQNLTVKLOLQALQNGSVLSEDSKRLNTILNTMTSTIYTGKVCNPDNPQEC-L 124
Db 93 DTVQSSDADIRROAEHLKLGASALNADDYLALQNAISSMQNTATATVCSYTNRSDCSL 152
Qy 125 LLEPGLNETMANSLDYNERLWAWESRVSQKLRPLYEYVVLKNEMARANHYEDYDGY 184
Db 153 TLEPHIQERLSRDPALAWYRWREHDKSGTPTMRQNFABYVRLTKASQLNGHRSYADY 212
Qy 185 WRGDYEVNGVDGYDSRGQLIEDVEHTFBEIKPLYEHLHAYVRACLNNAY-PSYISPIGC 243
Db 213 WQQFYE-----DPDFER-----QLDATPKQLPLYLQGLGYRFRLRQHYGVDVMPAEGN 262
Qy 244 LPAHLIGDMWGRFTWNLVSLTVPFGQKNIDVTDAMVDQAWDAQRIKFAEAKFVSVGLP 303
Db 263 IPISLILGNWGSNELLDFTPYPEKPFVDVKAEMEKQGYTVQKLFEIGDQFFQSLGMR 322
Qy 304 NMTQGFWNSMLTDPGNQKAVCHPTAWDLGK-GDPRILMCTKVTWDDFLTAHHEMGHIQ 362
Db 323 ALPPSPFWNLVSLTRPDQ-RQVVCASAWDFYQSDVRIKQKTEVDVSHYFVYVHHELGHQ 381
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Qy 363 YMAYAAQPFLLRNGANEGFHEAVGIMSLSAATPKHLKSGILLSPDFQEDNETEINFL 422
Db 382 YLYQEQPAPVYRGAPNPGFHEAVGDIYALSVMASAKHLKAIGLIE-NGRLDEKSRINQLF 440
Qy 423 KOALITVGLTPTMYLWKRMVVFKEIPKQDMKMKWEMKREIIVGVVEPVPHDETYCDP 482
Db 441 KQALSKIIVFLPGYAVDKRYAVAFRNELDESNQWCGFMQSEFGGVBPVPRTEKDPDP 500
Qy 483 ASLPHVNSDYSIRYTRTYQFOQFQALCOAAKEGP-----LHKCDISNSTEAGQKL 536
Db 501 PAKYHVSADVEYLRVFAAAHIFQFQFHKVLCRKAGQYAPNNSRLTLDNCDIFGSKAAGRSL 560
Qy 537 FNLRLGKSEPTLALENVVGAKNNVRPLLNYERPLFTWLKQDNKNSVFGVSHSTWSP 594
Db 561 SQFLSKGNSRHWKYLEBFTGETEMDPAALFEYFEPYQWLKQEB--NSRLGVLGMP 616

RESULT 12
AS7533
peptidyl-dipeptidase A (EC 3.4.15.1) 67k precursor - fruit fly (Drosophila melanogaster)
N:Alternate names: angiotensin-converting enzyme
C:Species: Drosophila melanogaster
C:Date: 08-Feb-1996 #sequence_revision 08-Feb-1996 #text_change 26-Feb-1998
C:Accession: AS7533
R:Corneil, M.J.; Williams, T.A.; Lamango, N.S.; Coates, D.; Corvol, P.; Soubrier, F.; Hot
J. Biol. Chem. 270, 13613-13619, 1995
A:Title: Cloning and expression of an evolutionary conserved single-domain angiotensin c
A:Reference number: AS7533; MUID:95293950; PMID:7775412
A:Accession: AS7533
A>Status: Preliminary
A:Molecule type: mRNA
A:Residues: 1-615 <COR>
A:Cross-references: UNIPARC:UPI0000175887; GB:U25344
C:Genetics:
A:Gene: FlyBase:Anc
A:Cross-references: FlyBase:PBgn0012037
C:Superfamily: mammalian peptidyl-dipeptidase A
C:Keywords: peptidyl dipeptide hydrolase

Query Match 31.6%; Score 1022; DB 2; Length 615;
Best Local Similarity 35.6%; Pred. No. 1.6e-65;
Matches 213; Conservative 118; Mismatches 245; Indels 22; Gaps 9;

Qy 4 EQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNNMAGDKWSAFLEKQSTLAQ 63
Db 22 EIQAKYELNENLKLAKRTNVETEAAWAGSNITDENKKQKEISAEALAKPMKEVASDIT 81
Qy 64 MYPLQEIQLTVKLOLQALQNGSVLSEDSKRLNTILNTMTSTIYTGKVCNPDNPQEC 123
Db 82 KFQWRSYQSEDLKROFKALTKLGYAALPEDDYAEILLDTLSAMBESNFAKVKVCYKDKSTK 141
Qy 124 -LLLEPGLNETMANSLDYNERLWAWESRVSQKLRPLYEYVVLKNEMARANHYEDY 182
Db 142 DLALDPETEEVVISKSRDHEELAYRREFYDKAGTAVRSQFERYVELNTYKAAKLNFSA 201
Qy 183 DYWRGDYEVNGVDGYDSRGQLIEDVEHTFBEIKPLYEHLHAYVRACLNNAY-PSYISPI 241
Db 202 EAWLDEYE-----DDTFEEQLEDI---FADIRPLLPAWPWLCAPLRKHYGDVAVSET 251
Qy 242 GCLPAHLIGDMWGRFTWNLVSLTVPFGQKNIDVTDAMVDQAWDAQRIKFAEAKFVSVG 301
Db 252 GPIMPILLGNMAQWMSIADIVSPFPEKPLVDVSAEMEKQAYTFLKMFQMGDDFFTSKN 311
Qy 302 LPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGK-GDPRILMCTKVTWDDFLTAHHEMGH 360
Db 312 LTKLPQDFWDSKILEKPDGDRDLVCHASAWDFYLTDDVRIKQCRTVQDQFFTVHHELG 371
Qy 361 IQYDMAYAAQPFLLRNGANEGFHEAVGIMSLSAATPKHLKSGILLSPDFQEDNETEIN 420
Db 372 IQYFLOYQHFPVIRGTGANGPGFHEAVGVDLSVSTPKHLKIGLIE-KYVRDDEARINQ 430
Qy 421 LLKQALTVGLTPTMYLWKRMVVFKEIPKQDMKMKWEMKREIIVGVVEPVPHDETYC 480
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Db 431 LFTALDKIVFLPFAFTMDKYRSLRGEVDKANWNCAPWKLRLDEYSGTEPPVVRSEKDF 490
QY 481 DPASLPHVSNDSYFIRYRTTLQFOFOEALC-QAAKH-----EGPLHKCDISNSTEAGQ 534
Db 491 DAPAKTHISADVEYLYSLVFIIOFYKSACIKAGQYDFDNVELPLDNCDIYGSARAGA 550
QY 535 KLFNMLRLGKSEPTWLTALENVVGAKNMVRPLINYEPLFTWLKQON--KNSFVGVNST 590
Db 551 AFHNLMSGASKPWPDALEAFNGERIMSGKATAEYFEPLRVWLEAENIKNNVHIGWT 608

RESULT 13
T15792
hypothetical protein C42D8.5 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 09-Jul-2004
C:Accession: T15792
R:Halloworth, K.
submitted to the EMBL Data Library, April 1996
A:Description: The sequence of C. elegans cosmid C42D8.
A:Reference number: Z18405
A:Accession: T15792
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-907 <HAL>
A:Cross-references: UNIPROT:Q18581; UNIPARC:UPI000004A55C; EMBL:U56966; NID:g1293844; PI
A:Experimental source: strain Bristol N2; clone C42D8
C:Genetics:
A:Gene: CESP:C42D8.5
A:Map position: X
A:Introns: 140/3; 170/3; 194/3; 300/2; 467/3; 551/2; 600/2; 697/3; 774/2; 851/3

Query Match 19.7%; Score 635.5; DB 2; Length 907;
Best Local Similarity 27.1%; Pred. No. 1.8e-37;
Matches 166; Conservative 131; Mismatches 277; Indels 39; Gaps 18;
QY 4 EEOAKTFLDKFNHEAEDLFYQSSLASWNTNTNITEENVQNNMAGDKWSAFLEKQOSTLAQ 63
Db 178 BEKLSWLAGYEAEAKVLRVALSGWRYFNDAFSLKALDAEAEVLTMTFVRSISMQAK 237
QY 64 MYPLOBIQNLTVKLQALQOQSSVLSEDKSKRLTILNTMTSTYSTGKVCNPNPQRC 123
Db 238 QEDMASVTDEKVMRQLGYVSFEQMSALAPSRADYSQAQALNRDSKOSTICDKVPPPC 297
QY 124 LLEPLGLNEIMANSDYNERLWAWESRSEVGKQRLPELYEYVLKNEMARANHYEDYGD 183
Db 298 ALQKIDMDSIFRNEKXASRLQHLWVSYVTAIAKS-KPSYNNIITISNEGAKLNGFANGGA 356
QY 184 YVRGDYEVNG-VDGYDSRGQIEDVEHTFEBIKPLYEHLHAYVRAKLMNAY--PSYISP 240
Db 357 MRSADFMSKVKHKAEP---DLNKQIDKIYSTIQPPYQLLHAYMRQLAGIYSNPVGLSK 413
QY 241 IGCPLPAHLGDMWGRFTWTLNLYSLTVPGQKPNIDVTAMDV---QAMDAQRIKFAEAKF 296
Db 414 DGPPIPAHLFGSLDGGDWSAHYEQTKEFEES--ETPEAMLSAFNTQNTYTKKMFVTAVRY 471
QY 297 FVSVGLPNMTQGFWNSMLTDPGNQKAVCHP-TAWDL-GKGFRLMCTKVTMDDFLTA 354
Db 472 FKSAGPPLPKYSYTSISIFARWNS-KDMICHFAAALDMRAPNDPRVYKACAOGLGEPDFEOA 530
QY 355 HHMGHIOYDMAVAAQPFLLRNGAEGFHEAVGEIMSLSAATPKHLKSLGILLSPPQFQEDN 414
Db 531 HSLLVQTYQYLYKQDLSLFRQASPVITDANAFHLSTNPHLYLSQKLVPSSEHLDIK 590
QY 415 ETE-INFLKQALTIIVGTLPTFTYMLEKRWVFKGEIPKQDMKKWEMKREIVGVVEFV 473
Db 591 DSVIINKLYKESLESPTKLPFTIAADNRWYELFDGTVPKNKLNDRWWEIRNKYEGVRSPQ 650
QY 474 PHDETYCDPASLEH-VSNDYSFIRYTTTL-----YQFOFOALCOAA---KHEGPLHKC 524
Db 651 PYNTSLND--ALIHNSVSQVHS---PATRTLISYVLKFOILKALCQRELFWLSEG----C 701

QY 525 DISNSTEAGQKLFNNRLGKSEPTWLTALENVVGAKNMVRPLINYEPLFTWLKQONK-- 582
Db 702 ILSEDTT--EKLRETMKLGSSITWLKALEMISGKELDAQPLLEYEPLINLWLRNTNEID 759
QY 583 NSFVGVNSTDWSPY 595
Db 760 QVVVGWMDGEGTTP 772
RESULT 14
C83696
hypothetical protein BH0371 [imported] - Bacillus halodurans (strain C-125)
C:Species: Bacillus halodurans
C:Date: 01-Dec-2000 #sequence_revision 01-Dec-2000 #text_change 09-Jul-2004
C:Accession: C83696
R:Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Maui, N.; Fujii, F.; Hiran
Nucleic Acids Res. 28, 4317-4331, 2000
A:Title: Complete genome sequence of the alkaliphilic bacterium Bacillus halodurans and i
A:Reference number: A83650; MUID:20512582; PMID:11058132
A:Accession: C83696
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-532 <STO>
A:Cross-references: UNIPROT:Q9KFV0; UNIPARC:UPI00000C3863; GB:AP001508; GB:BA000004; NID:
A:Experimental source: strain C-125
C:Genetics:
A:Gene: BH0371

Query Match 4.9%; Score 157; DB 2; Length 532;
Best Local Similarity 21.1%; Pred. No. 0.002;
Matches 118; Conservative 83; Mismatches 213; Indels 144; Gaps 29;
QY 4 EEOAKTFLDKFNHEAEDLFYQSSLASWNTNTNITEENVQNNMAGDKWSAFLEKQES--- 59
Db 3 EODIERFLSEQNKRVEDLYQPVLLNHNWVATTGEOEWSKHEQSUSEYWAHFSRSEFOK 62
QY 60 -----TLAQMYPQLQEIQNLTVKLQALQOQSSVLSEDKSKRLTILNTMTSTIY 109
Db 63 VTRFRKIDSLPLMQRRQLDDLHDKMKIQPE--EGTRQOILSLE--KKISHVFTTFQPV 118
QY 110 STGKVCNPDNPQECLELLPEGLNEIMANSLDYNERLWAWESRSEVGKQRLPELYEYVLK 169
Db 119 NGSRYSNNE-----LLDILRYDLDERHRRKQAWFA-SKEVGKRTKDKDLLQIRKR 166
QY 170 NEMARANHYEDGYVRGDYEVNGVDGYDSRGQLIEDVEHT---FEBIKPLYEHLHAYV 226
Db 167 NEVARNLGFETP-----YHMSFAQQLDLEQTFAMFETIKKSSDOAFRMI 211
QY 227 -----RAKLMNAYPSYISPIGCLPAHLGDMWGRFTWTLNLYSLTVPGQK-PNIDVTDA 278
Db 212 KDEIDEBRAKVLKIKKODLRP-----WDYVDPFFQEAPSIHVD- 250
QY 279 MVDQAWDAQRIFKEAEKFFVSVGLPNMTQGFWNSMLTDPGNVQK-AVCHPTAMDGLKGD 337
Db 251 -FDSFYKQDLEQVVSQTFQAMELP--IDDILKRSGLYPKRNQNPFGFC--TDMD-RRGD 304
QY 338 FRILMCTKVTMDDFLTAHHENGH-IQYDMAVAAQPFLLRNGAEGFHEAVGEIMSLSAAT 396
Db 305 IRVLLNLDQSMYWTALLHFEFGHAVYFKFIDSRPFLLR-----FHT-----SHTLTT 351
QY 397 PKHLKSLGILLS--PDFQES-----DNET-----EINFLLKQALTIIVGTLPTFTYMLEKRW 443
Db 352 EASALFFGRTMKAWEYERFLGIDRETCTERIGNMEKMLQRM-VVST-----RW 400
QY 444 MV-----FKG-----EIPKQDMKKWEMKREIVGVVEPVPDDETYCDPASLFHVS----- 490
Db 401 MLTFSFEKSLYEDPDQDINALWKLVEIKYQMAP--PEDTGSPPDMAAKMHFSLAPVYQ 458
QY 491 DY-----SFIRYVYTRT 501
Db 459 DYLLGEMAASQLHHYIKT 476

Search completed: March 28, 2006, 11:17:18
Job time : 24.8626 secs

RESULT 15

AF1310
probable thermostable carboxypeptidases homolog lmo1886 [imported] - Listeria monocytogenes
C:Species: Listeria monocytogenes
C:Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 05-Oct-2004
C:Accession: AF1310
R:Glaser, P.; Prangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloeker, H.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Feihl, H.H.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A:Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkat, G.; Madueno, E.; Maitournam, A.; Maier, C.; Schluter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehlend, O., C.; Schlueter, J.; Simeos, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehlend, O.
A:Title: Comparative genomics of Listeria species.
A:Reference number: AB1077; MUID:21537279; PMID:11679669
A:Accession: AF1310
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-502 <GLA>
A:Cross-references: UNIPROT:Q8Y616; UNIPARC:UPI00000CF1B7; GB:NC_003210; PIDN:CAC99964.1
A:Experimental source: strain EGD-e
C:Genetics:
A:Gene: lmo1886
C:Superfamily: Zn-dependent carboxypeptidase, Taq type

Query Match 4.8%; Score 154; DB 2; Length 502;
Best Local Similarity 20.1%; Pred. No. 0.003;
Matches 128; Conservative 101; Mismatches 215; Indels 192; Gaps 35;

Qy 2 TIEEQATFLDKFNHEADLIFYOSSLASWNYT-----NITEENVQNNA 47
Db ||| :
4 TLEEFLAYIKMEALEAL-----ALVMDLRGTAPAKMGEGRSDIVGLSEIFNQTS 59
Qy :
48 GDKWSAFLEKQSTLAQMYPLEIQNTLVQLQALQQNGSSVLSDSKREL-----N 99
Db :
60 -EEMAAFIAGLN-----QDENLS-EITRKILE--SQTDLNKPKPKEVAEYT 106
Qy TLNTMTSYITGKVCNPDPQBCLLEPLGLEINMANSLDYNERLMAWESWRSEVGKQLR 159
Db ||| :
107 KLVAQAETAWTTAREQN-----DFAAPEFLTIL----- 136
Qy 160 PLVEEYVVVKNEMARANHYEDGYNRGRDYEVNGVDGY--DYSRGQLIEDVHTPEIKP 217
Db ||| :
137 -----EMKR-----KFVEYW--GYEENKYDTLLDQYEFGVTVSVIDSVFEKVR- 177
Qy 218 LYEHLYAVRAKLNNAYPSVISPTGCLPAHLGLDMGMRFTNLYSLTVPFGQKPNDIVTD 277
Db ||| :
178 --DGIMA-IREKIENE-----GVKPDATILN 200
Qy 278 AMVDQAWDAQRIFKEARKFFVSVGLPNMTGGFWNSMLTDPGNVQKAVCHPTAMDLGKGD 337
Db ||| :
201 TKISEA-----KQKEFSIRI-LNKWGDFP-----EAGRLDET-VHPFATGLNTGD 243
Qy 338 FRILMCCTKVTMDPLTA-----HHEMGHIQV----DMAYAAQPFLLRNEGPHAEVG-- 387
Db ||| :
244 VRI--TTRYENDFKMAVFGTIEHGHAIVEQNFDALVGTGP--LANGSMGHESQSLE 299
Qy 388 -EIM--SLSAATPKHLKSIGLLSPDQF---BD-----NETEIFLLKQALTIVGTLP- 433
Db ||| :
300 YEIIIGSSLAPFKSNVADFQAITKPAFDQVKLEDYFRVAVNISSSLIREEADTL--TYPL 357
Qy 434 ---FTYMLEKRWNVFKGEIPKQWMKKWEMKREIVGVVEPVPHDETCDPASLFHVN 490
Db ||| :
358 HIMIRYEELEK---ALINGELEVKDLPKAWGDKYEEYGI---RPDNDTNGVLQDIHWAG 411
Qy 491 DYSFIRYYRTLL-YQOFQEBALCOAAKHGPHKCDISNSTEAGOKLF---NMLRGKSE 546
Db ||| :
412 DFGYFPSPYALGLMYAAQPFNQW----QKEIPNIDAIASDDYSELKIWLTEHVHKGKTK 467
Qy 547 PWTLLAENVVGAKNMNVRPLLANYPELFTWLKDQNX 582
Db KPTEILDITDTTG-EGINPTYLLDLLLEKRYAYVYQFNK 502

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GenCore version 5.1.7
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OM protein - protein search, using sw model

Run on: March 28, 2006, 11:03:07 ; Search time 124.785 Seconds
(without alignments)
3364.096 Million cell updates/sec

Title: US-10-659-000-4
Perfect score: 3231
Sequence: 1 STIEGQAKTFLDKFHEAED.....WLKDQNKNSFVGWSTDNPSY 595

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 2166443 seqs, 705528306 residues

Total number of hits satisfying chosen parameters: 2166443

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Uniprot 05.80.*

1: uniprot_sprot.*

2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3231	100.0	805	1	ACE2_HUMAN
2	3193	98.8	805	1	ACE2_PONPY
3	2823	87.4	805	1	ACE2_FELCA
4	2794	86.5	805	1	ACE2_PAGLA
5	2757	85.3	805	1	ACE2_RAT
6	2755	85.3	805	1	ACE2_MOUSE
7	2710.5	83.9	804	1	ACE2_BOVIN
8	2018	62.5	785	2	Q5U380_BRARE
9	1613	49.9	652	2	Q4SHR0_TETNG
10	1345	41.6	1314	1	ACE_MOUSE
11	1335	41.3	732	1	ACE_T_HUMAN
12	1335	41.3	739	2	Q8N710_HUMAN
13	1335	41.3	1306	1	ACE_HUMAN
14	1334	41.3	732	1	ACET_MOUSE
15	1334	41.3	1015	2	Q8K233_MOUSE
16	1334	41.3	1187	2	Q5XK22_MOUSE
17	1334	41.3	1312	1	ACE_MOUSE
18	1328	41.1	1144	2	Q4S1M8_TETNG
19	1327	41.1	732	1	ACET_PANTR
20	1327	41.1	1304	1	ACE_PANTR
21	1310	40.5	775	1	ACET_RAT
22	1310	40.5	1313	1	ACE_RAT
23	1307	40.5	1193	1	ACE_CHICK
24	1301	40.3	694	2	Q15540_HUMAN
25	1295	40.1	724	2	Q4W1E4_SHEEP
26	1281	39.6	737	1	ACET_RABIT
27	1281	39.6	1310	1	ACE_RABIT
28	1236	38.3	616	1	ACE_THETS
29	1176.5	36.4	625	2	Q6XK62_LOCMI
30	1139.5	35.3	627	2	Q5WPT4_LUTIO
31	1092.5	33.8	619	2	Q8BE93_SHEON

32	1088	33.7	617	2	Q7PM22_ANOGA
33	1078	33.4	648	2	Q9NDS8_BOMMO
34	1068	33.1	615	1	ACE_DROME
35	1054.5	32.6	611	1	ACE_HAEIE
36	1054	32.6	660	2	Q17248_BOOMI
37	1025.5	31.7	630	1	ACER_DROME
38	1005.5	31.1	614	2	QANKG3_9DELIT
39	984.5	30.5	631	2	Q7NGM5_GLOVI
40	965	29.9	672	2	Q4URZ8_XANCP
41	965	29.9	672	2	Q8PBK3_XANCP
42	964	29.8	672	2	Q8PN56_XANAC
43	961	29.7	977	2	Q7O9W7_ANOGA
44	949	29.4	686	2	Q5GWX8_XANOR
45	943.5	29.2	567	2	Q5VLH5_XANOR

ALIGNMENTS

RESULT 1
ACE2_HUMAN
ID ACE2_HUMAN STANDARD; PRT; 805 AA.
AC Q9BYF1; Q6UWPO; Q86WT0; Q9NRA7; Q9UFZ6;
DT 13-SEP-2005 (Rel. 48, Created)
DT 13-SEP-2005 (Rel. 48, Last sequence update)
DT 13-SEP-2005 (Rel. 48, Last annotation update)
DE Angiotensin-converting enzyme 2 precursor (SC 3.4.17.-) (ACE-related carboxypeptidase) (Angiotensin-converting enzyme homolog) (ACEH).
GN Name=ACE2; ORFNames=UNQ868/PRO1885;
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae;
OC Homo.
OX NCBI_TaxID=9606;
RN [1]
RP NUCLEOTIDE SEQUENCE [MRNA] (ISOFORM 1), TISSUE SPECIFICITY, FUNCTION, AND ENZYME REGULATION.
RC TISSUE=Heart;
RX MEDLINE=20429895; PubMed=10969042;
RA Donoghue M., Hsieh F., Baronas E., Godbout K., Gosselin M., Stagliano N., Donovan M., Woolf B., Robison K., Jeyaseelan R., Breitbart R.E., Acton S.;
RA "A novel angiotensin-converting enzyme-related carboxypeptidase (ACE2) converts angiotensin I to angiotensin 1-9.";
Circ. Res. 87:E1-E9(2000).
[2]
RP NUCLEOTIDE SEQUENCE [MRNA] (ISOFORM 1), TISSUE SPECIFICITY, NUCLEOTIDE SEQUENCE [MRNA] (ISOFORM 1), TISSUE SPECIFICITY, FUNCTION, AND ENZYME REGULATION.
RC TISSUE=Lymphoma;
RX MEDLINE=20517872; PubMed=10924499; DOI=10.1074/jbc.M002615200;
RA Tipnis S.R., Hooper N.M., Hyde R., Karran E., Christie G., Turner A.J.;
RA "A human homolog of angiotensin-converting enzyme. Cloning and functional expression as a captopril-insensitive carboxypeptidase.";
J. Biol. Chem. 275:33238-33243(2000).
[3]
RP NUCLEOTIDE SEQUENCE [MRNA] (ISOFORM 1), TISSUE SPECIFICITY, SUBCELLULAR LOCATION, AND ENZYME REGULATION.
RC TISSUE=Testis;
RX PubMed=15231706; DOI=10.1210/en.2004-0443;
RA Douglas G.C., O'Bryan M.K., Hedger M.P., Lee D.K.L., Yarski M.A., Smith A.I., Lew R.A.;
RA "The novel angiotensin-converting enzyme (ACE) homolog, ACE2, is selectively expressed by adult Leydig cells of the testis.";
Endocrinology 145:4703-4711(2004).
[4]
RN NUCLEOTIDE SEQUENCE [MRNA] (ISOFORM 1), AND VARIANT SER-638.
RP TISSUE=Lung, and Testis;
RX PubMed=15937940; DOI=10.1002/ajmg.a.30779;
RA Itoyama S., Keicho N., Hijikata M., Quy T., Phi N.C., Long H.T., Ha I.D., Ban V.V., Matsuhashita I., Yanai H., Kirikae F., Kirikae T., Kuratsuji T., Sasazuki T.;
RA "Identification of an alternative 5'-untranslated exon and new

RT polymorphisms of angiotensin-converting enzyme 2 gene: Lack of
RT association with SARS in the Vietnamese population.;"
RL Am. J. Med. Genet. 136:52-57(2005).
RN [5]
RP NUCLEOTIDE SEQUENCE [MRNA] (ISOFORM 1).
RA Suzuki Y., Watanabe M., Sugano S.;
RT "Cloning, expression analysis and chromosomal localization of a novel
RT ACE like enzyme.;"
RL Submitted (JUL-2000) to the EMBL/GenBank/DBJ databases.
RN [6]
RP NUCLEOTIDE SEQUENCE [LARGE SCALE MRNA] (ISOFORM 2).
RX MEDLINE=22887296; PubMed=12975309; DOI=10.1101/gr.1293003;
RA Clark H.F., Gurney A.L., Abaya E., Baker K., Baldwin D.T., Brush J.,
RA Chen J., Chow B., Chui C., Crowley C., Currell B., Deuel B., Dowd P.,
RA Eaton D., Foster J.S., Grimaldi C., Gu Q., Hass P.E., Heldens S.,
RA Huang A., Kim H.S., Klimowski L., Jin Y., Johnson S., Lee J.,
RA Lewis L., Liao D., Mark M.R., Robbie E., Sanchez C., Schoenfeld J.,
RA Seshagiri S., Simmons L., Singh J., Smith V., Stinson J., Vagts A.,
RA Vandlen R.L., Watanabe C., Wiedand D., Woods K., Xie M.-H.,
RA Yanagura D.G., Yi S., Yu G., Yuan J., Zhang M., Zhang Z., Goddard A.D.,
RA Wood W.I., Godowski P.J., Gray A.M.;
RT "The secreted protein discovery initiative (SPDI), a large-scale
RT effort to identify novel human secreted and transmembrane proteins: a
RT bioinformatics assessment.;"
RL Genome Res. 13:2265-2270(2003).
RN [7]
RP NUCLEOTIDE SEQUENCE [GENOMIC DNA] (ISOFORM 1), AND VARIANT ARG-26.
RA Rieder M.J., Carrington D.P., da Ponte S.H., Hastings N.C.,
RA Ahearn M.O., Kuldane S.A., Rajkumar N., Toth E.J., Yi Q.,
RA Nickerson D.A.;
RT "SeattlesNP. NHLBI HL66682 program for genomic applications, UW-
RT FHCR, Seattle, WA (URL: <http://pga.gs.washington.edu>).;"
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
RN [8]
RP NUCLEOTIDE SEQUENCE [LARGE SCALE MRNA] (ISOFORM 1).
RT TISSUE=Brain, and Testis;
RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Donald M.F., Casavant T.L., Schestz T.E.,
RA Brownstein M.J., Ussid T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettelman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakeley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywiniski M.I., Skalska U., Smalhus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences.;"
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [9]
RP NUCLEOTIDE SEQUENCE [LARGE SCALE MRNA] OF 2-805 (ISOFORM 1).
RT TISSUE=Testis;
RG The German cDNA consortium;
RL Submitted (AUG-1999) to the EMBL/GenBank/DBJ databases.
RN [10]
RP PROTEIN SEQUENCE OF 679-689, IDENTIFICATION BY MASS SPECTROMETRY, AND
RP INTERACTION WITH ITGB1.
RX PubMed=15276642; DOI=10.1016/j.bbdis.2004.05.005;
RA Lin Q., Keller R.S., Weaver B., Zigman L.S.;
RT "Interaction of ACE2 and integrin beta1 in failing human heart.;"
RL Biochim. Biophys. Acta 1689:175-178(2004).
RN [11]
RP TISSUE SPECIFICITY.
RX MEDLINE=22347248; PubMed=12459472;
RA Harmer D., Gilbert M., Borman R., Clark K.L.;

RT "Quantitative mRNA expression profiling of ACE 2, a novel homologue of
RT angiotensin converting enzyme.;"
RL FEBS Lett. 532:107-110(2002).
RN [12]
RP BIOPHYSICOCHEMICAL PROPERTIES, ENZYME REGULATION, AND COFACTOR.
RX PubMed=11815627; DOI=10.1074/jbc.M200581200;
RA Vickers C., Hales P., Kaushik V., Dick L., Gavin J., Tang J.,
RA Godbout K., Parsons T., Baronas E., Hsieh F., Acton S., Patane M.A.,
RA Nichols A., Tummino P.;
RT "Hydrolysis of biological peptides by human angiotensin-converting
RT enzyme-related carboxypeptidase.;"
RL J. Biol. Chem. 277:14838-14843(2002).
RN [13]
RP FUNCTION, INTERACTION WITH HCOV-SARS S PROTEIN, GLYCOSYLATION, AND
RP IDENTIFICATION BY MASS SPECTROMETRY.
RX PubMed=14647384; DOI=10.1038/nature02145;
RA Li W., Moore M.J., Vasilieva N., Sui J., Wong S.-K., Berne M.A.,
RA Somasundaran M., Sullivan J.L., Iuzriaga K., Greenough T.C., Choe H.,
RA Farzan M.;
RT "Angiotensin-converting enzyme 2 is a functional receptor for the SARS
RT coronavirus.;"
RL Nature 426:450-454(2003).
RN [14]
RP INDUCTION.
RX PubMed=15151696; DOI=10.1186/1741-7015-2-19;
RA Goulter A.B., Goddard M.J., Allen J.C., Clark K.L.;
RT "ACE2 gene expression is up-regulated in the human failing heart.;"
RL BMC Med. 2:19-19(2004).
RN [15]
RP TISSUE SPECIFICITY.
RX PubMed=15141377; DOI=10.1002/path.1570;
RA Hamming I., Timens W., Bulthuis M.L.C., Lely A.T., Navis G.J.,
RA van Goor H.;
RT "Tissue distribution of ACE2 protein, the functional receptor for SARS
RT coronavirus. A first step in understanding SARS pathogenesis.;"
RL J. Pathol. 203:631-637(2004).
RN [16]
RP INTERACTION WITH HCOV-SARS S PROTEIN.
RX PubMed=15452268; DOI=10.1128/JVI.78.20.11429-11433.2004;
RA Li W., Greenough T.C., Moore M.J., Vasilieva N., Somasundaran M.,
RA Sullivan J.L., Farzan M., Choe H.;
RT "Efficient replication of severe acute respiratory syndrome
RT coronavirus in mouse cells is limited by murine angiotensin-converting
RT enzyme 2.;"
RL J. Virol. 78:11429-11433(2004).
RN [17]
RP TISSUE SPECIFICITY, AND INDUCTION.
RX PubMed=15671045; DOI=10.1093/eurheartj/ehi114;
RA Burrell L.M., Risvanis J., Kubota E., Dean R.G., MacDonald P.S.,
RA Lu S., Tikellis C., Grant S.L., Lew R.A., Smith A.I., Cooper M.E.,
RA Johnston C.I.;
RT "Myocardial infarction increases ACE2 expression in rat and humans.;"
RL Eur. Heart J. 26:369-375(2005).
RN [18]
RP INTERACTION WITH HCOV-SARS S PROTEIN, AND MUTAGENESIS.
RX PubMed=15791205; DOI=10.1038/sj.emboj.7600640;
RA Li W., Zhang C., Sui J., Kuhn J.H., Moore M.J., Luo S., Wong S.-K.,
RA Huang I.-C., Xu K., Vasilieva N., Murakami A., He Y., Marasco W.A.,
RA Guan Y., Choe H., Farzan M.;
RT "Receptor and viral determinants of SARS-coronavirus adaptation to
RT human ACE2.;"
RL EMBO J. 24:1634-1643(2005).
RN [19]
RP PROTEOLYTIC CLEAVAGE.
RX PubMed=15983030; DOI=10.1074/jbc.M505111200;
RA Lambert D.W., Yarski M.A., Warner F.J., Thornhill P., Parkin E.T.,
RA Smith A.I., Hooper N.M., Turner A.J.;
RT "Tumor necrosis factor-alpha convertase (ADAM17) mediates regulated
RT ectodomain shedding of the SARS-CoV receptor, angiotensin-converting
RT enzyme-2 (ACE2).;"
RL J. Biol. Chem. 280:0-0(2005).
RN [20]
RP INTERACTION WITH HCOV-NL63 S PROTEIN.


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RX PubMed=15897467; DOI=10.1073/pnas.0409465102;
RA Hofmann H., Pyrc K., van der Hoek L., Geier M., Berkhout B.,

Query Match      100.0%; Score 3231; DB 1; Length 805;
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Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 61 LAQMYPLQEIQNLTVKLOLQALQNGSSVLSEDKSKRLNTILNTMTSTYSTGKVCNPNP 120
DB 79 LAQMYPLQEIQNLTVKLOLQALQNGSSVLSEDKSKRLNTILNTMTSTYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSVGKQLRPLYEEYVVLKKNEMARANHED 180
DB 139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSVGKQLRPLYEEYVVLKKNEMARANHED 198
QY 181 YGDYWRGDEYVNGVDYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
DB 199 YGDYWRGDEYVNGVDYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
QY 241 IGCPLPAHLGDMWGRFNTNLSYTVFPQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSU 300
DB 259 IGCPLPAHLGDMWGRFNTNLSYTVFPQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSU 318
QY 301 GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGDPRILMCTKYTMDDFLTAAHEMGGH 360
DB 319 GLPNMTQGFWENSMLTDPGNVQKAVCHPTAWDLGKGDPRILMCTKYTMDDFLTAAHEMGGH 378
QY 361 IQYDMAYAAQPLLRNGANEGFHEAVGEIMSLSAATPKHLKSGILSLSPQSDNETEINF 420
DB 379 IQYDMAYAAQPLLRNGANEGFHEAVGEIMSLSAATPKHLKSGILSLSPQSDNETEINF 438
QY 421 LKQALTIYVGTLPFTYMLEKRWMMVKGEIPKQDMKKWEMKREIVGVVEPVPHDETTC 480
DB 439 LKQALTIYVGTLPFTYMLEKRWMMVKGEIPKQDMKKWEMKREIVGVVEPVPHDETTC 498
QY 481 DPASLPHVSNDSYFIRYRTTLYQFQEQALCOAAKHEGHLKHKCDISNTEAGQKLFNNML 540
DB 499 DPASLPHVSNDSYFIRYRTTLYQFQEQALCOAAKHEGHLKHKCDISNTEAGQKLFNNML 558
QY 541 RLKGSBPWTALENVVGAKNMVRPLLNTFPEPLFTWLKQNKNSFVGWSTDSWSPY 595
DB 559 RLKGSBPWTALENVVGAKNMVRPLLNTFPEPLFTWLKQNKNSFVGWSTDSWSPY 613

RESULT 2
ACE2_PONPY STANDARD; PRT; 805 AA.
AC QSRFN1;
DT 13-SEP-2005 (Rel. 48, Created)
DT 13-SEP-2005 (Rel. 48, Last sequence update)
DT 13-SEP-2005 (Rel. 48, Last annotation update)
DE Angiotensin-converting enzyme 2 precursor (EC 3.4.17.-) (ACE-related
DE carboxypeptidase).
GN Name=ACE2;
OS Pongo pygmaeus (Orangutan).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae;
OC Pongo.
OC NCBI_TaxID=9600;
RN [1]
RP NUCLEOTIDE SEQUENCE [LARGE SCALE MRNA].
RC TISSUE=kidney;
RG The German cDNA consortium;
RL Submitted (NOV-2004) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Carboxypeptidase which converts angiotensin I to
CC angiotensin 1-9, a peptide of unknown function, and angiotensin II
CC to angiotensin 1-7, a vasodilator. Also able to hydrolyze apelin-
CC 13 and dynorphin-13 with high efficiency. May be an important
CC regulator of heart function (By similarity).

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CC -!- COPACTOR: Binds 1 zinc ion per subunit (By similarity).
CC -!- COPACTOR: Binds 1 chloride ion per subunit (By similarity).
CC -!- SUBCELLULAR LOCATION: Type I membrane protein. Processing by
CC ADAM17 may lead to a secreted protein (By similarity).
CC -!- SIMILARITY: Belongs to the peptidase M2 family.
CC -----
CC This Swiss-Prot entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use as long as its content is in no way modified and this statement is not
CC removed.
CC -----
CC EMBL; CR857122; CAH89426.1; -; mRNA.
DR SMR; QSRFN1; 19-615.
DR InterPro; IPR006025; Pept_M_Zn_BS.
DR InterPro; IPR001548; Peptidase_M2.
DR Pfam; PF01401; Peptidase_M2; 1.
DR PRINTS; PR00791; PEPDPTASEA.
DR ProDom; PD004184; Peptidase_M2; 1.
DR PROSITE; PS00142; ZINC_PROTEASE; 1.
KW Carboxypeptidase; Chloride; Glycoprotein; Hydrolase; Metal-binding;
KW Metalloprotease; Protease; Signal; Transmembrane; Zinc.
FT SIGNAL 1 17
FT CHAIN 18 805 Angiotensin-converting enzyme 2.
FT TOPO_DOM 18 740 Extracellular (Potential).
FT TRANSMEM 741 761 Potential.
FT TOPO_DOM 762 805 Cytoplasmic (Potential).
FT ACT_SITE 375 375 By similarity.
FT ACT_SITE 505 505 By similarity.
FT METAL 374 374 Zinc (catalytic) (By similarity).
FT METAL 378 378 Zinc (catalytic) (By similarity).
FT METAL 402 402 Zinc (catalytic) (By similarity).
FT BINDING 169 169 Chloride (By similarity).
FT BINDING 273 273 Substrate (By similarity).
FT BINDING 345 345 Substrate (By similarity).
FT BINDING 346 346 Substrate (By similarity).
FT BINDING 371 371 Substrate (By similarity).
FT BINDING 477 477 Chloride (By similarity).
FT BINDING 481 481 Chloride (By similarity).
FT BINDING 515 515 Substrate (By similarity).
FT CARBOHYD 53 53 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 90 90 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 103 103 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 322 322 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 432 432 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 546 546 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 690 690 N-linked (GlcNAc...) (Potential).
FT DISULFID 133 141 By similarity.
FT DISULFID 344 361 By similarity.
FT DISULFID 530 542 By similarity.
SQ SEQUENCE 805 AA; 92605 MW; 9F68CC1ACBC763F1 CRC64;

Query Match      98.8%; Score 3193; DB 1; Length 805;
Best Local Similarity 98.8%; Pred. No. 2.1e-217;
Matches 588; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 1 STIEBQAKTFLDKFNHEADLFYQSSLASWNYNTNTEENVQNMNAGDKWSAFLKEQST 60
DB 19 STIEBQAKTFLDKFNHEADLFYQSSLASWNYNTNTEENVQNMNAGDKWSAFLKEQST 78
QY 61 LAQMYPLQEIQNLTVKLOLQALQNGSSVLSEDKSKRLNTILNTMTSTYSTGKVCNPNP 120
DB 79 LAQMYPLQEIQNLTVKLOLQALQNGSSVLSEDKSKRLNTILNTMTSTYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSVGKQLRPLYEEYVVLKKNEMARANHED 180
DB 139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSVGKQLRPLYEEYVVLKKNEMARANHED 198
QY 181 YGDYWRGDEYVNGVDYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
DB 199 YGDYWRGDEYVNGVDYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
QY 241 IGCPLPAHLGDMWGRFNTNLSYTVFPQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSU 300

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259 IGCLPAHLGLDMWGRFTWNTLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSU 318
QY 301 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVWDDFLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNVOKVCHPTAWDLGKGFRLMCTKVWDDFLTAHHEMGH 378
QY 361 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
Db 379 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438
QY 421 LLKQALTIIVGTLPTFTYMLEKRWVFKGEIPKQNMKKWEMKREIVGVVPEVPHDETYC 480
Db 439 LLKQALTIIVGTLPTFTYMLEKRWVFKGEIPKQNMKKWEMKREIVGVVPEVPHDETYC 498
QY 481 DPASLPHVSNDSYFRYYRTLYQFOFQALCOAAKHEGPLHKCDISNSTEAGOKLFNML 540
Db 499 DPASLPHVSNDSYFRYYRTLYQFOFQALCOAAKHEGPLHKCDISNSTEAGOKLFNML 558
QY 541 RLKSEPTWLTALENVVGAKNMVRPLNLYFPELFTWLKQDNKNSFVGWSTWDSY 595
Db 559 RLKSEPTWLTALENVVGAKNMVRPLNLYFPELFTWLKQDNKNSFVGWSTWDSY 613

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RESULT 3

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ACE2_FELCA
ID ACE2_FELCA STANDARD; PRT; 805 AA.
AC Q56H28;
DT 13-SEP-2005 (Rel. 48, Created)
DT 13-SEP-2005 (Rel. 48, Last sequence update)
DT 13-SEP-2005 (Rel. 48, Last annotation update)
DE Angiotensin-converting enzyme 2 precursor (EC 3.4.17.-) (ACE-related
DE carboxypeptidase).
GN Name=ACE2;
OS Felis silvestris catus (Cat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Laurasiatheria; Carnivora; Fissipedia; Felidae;
OC Felinae; Felis.
OX NCBI_TaxID=9685;
RN [1]
RP NUCLEOTIDE SEQUENCE [MRNA].
RA Wang C., Guo A.Z., Chen H.C.;
RT "Identification of cat ACE2 gene and its potential function as a SARS-
RT Cov receptor.";
RL Submitted (MAR-2005) to the EMBL/GenBank/DBAJ databases.
CC -!- FUNCTION: Carboxypeptidase which converts angiotensin I to
CC angiotensin 1-9, a peptide of unknown function, and angiotensin II
CC to angiotensin 1-7, a vasodilator. Also able to hydrolyze apelin-
CC 13 and dynorphin-13 with high efficiency. May be an important
CC regulator of heart function (By similarity).
CC -!- COFACTOR: Binds 1 zinc ion per subunit (By similarity).
CC -!- COFACTOR: Binds 1 chloride ion per subunit (By similarity).
CC -!- SUBCELLULAR LOCATION: Type I membrane protein. Processing by
CC ADAM17 may lead to a secreted protein (By similarity).
CC -!- SIMILARITY: Belongs to the peptidase M2 family.
CC -----
CC This Swiss-Prot entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use as long as its content is in no way modified and this statement is not
CC removed.
CC -----
CC EMBL; AY957464; AAX59005.1; -; mRNA.
CC InterPro; IPR006025; Pept_M_zn_BS.
CC InterPro; IPR001548; Peptidase_M2.
CC Pfam; PF01401; Peptidase_M2; 1.
CC PRINTS; PR00791; PEPTIDTASEA.
CC PROSITE; PS00142; ZINC_PROTEASE; 1.
CC Carboxypeptidase; Chloride; Glycoprotein; Hydrolase; Metal-binding;
KW Metalloprotease; Protease; Signal; Transmembrane; zinc.
FT SIGNAL 1 17 Potential.
FT CHAIN 18 805 Angiotensin-converting enzyme 2.
FT TOPO_DOM 18 740 Extracellular (Potential).

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TRANSMEM 741 761 Potential.
FT TOPO_DOM 762 805 Cytoplasmic (Potential).
FT ACT_SITE 375 375 By similarity.
FT ACT_SITE 375 375 By similarity.
FT METAL 374 374 Zinc (catalytic) (By similarity).
FT METAL 378 378 Zinc (catalytic) (By similarity).
FT METAL 402 402 Zinc (catalytic) (By similarity).
FT BINDING 169 169 Chloride (By similarity).
FT BINDING 273 273 Substrate (By similarity).
FT BINDING 345 345 Substrate (By similarity).
FT BINDING 346 346 Substrate (By similarity).
FT BINDING 371 371 Substrate (By similarity).
FT BINDING 477 477 Chloride (By similarity).
FT BINDING 481 481 Chloride (By similarity).
FT BINDING 515 515 Substrate (By similarity).
FT CARBOHYD 53 53 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 90 90 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 216 216 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 299 299 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 322 322 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 546 546 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 660 660 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 690 690 N-linked (GlcNAc...) (Potential).
FT DISULFID 133 141 By similarity.
FT DISULFID 344 361 By similarity.
FT DISULFID 530 542 By similarity.
SQ SEQUENCE 805 AA; 92708 MW; 9F41A2EF300BE19E CRC64;

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Query Match 87.4%; Score 2823; DB 1; Length 805;
Best Local Similarity 86.1%; Pred. No. 3.6e-191;
Matches 512; Conservative 37; Mismatches 46; Indels 0; Gaps 0;
QY 1 STTSEQAATFLDKFNHAEADLFYQSSLSASWNYNTNITEENVQNNMAGDKWSAFLEKQST 60
Db 19 STTSELAATFLKFNHAEADLFYQSSLSASWNYNTNITEENVQNNMAGDKWSAFLEKQST 78
QY 61 LAQMPLOEIQNLVTKVQLQALQONGSSVLSSEKRLNTILNTWSTIYSTGKVCNPNP 120
Db 79 LAKTYPLAEIHNTTVKRLQALQONGSSVLSSEKRLNTILNTWSTIYSTGKVCNPNP 138
QY 121 QECILLEPGLNEIMANSLDYNLERLWAMESVSGKQLRPLYEYVVLKNMAMRANHYED 180
Db 139 QECILLEPGLDDIMENSKDYNLERLWAMESVSGKQLRPLYEYVVLKNMAMRANHYED 198
QY 181 YGDYWRGDIYVNGVDYDSRGQIEDVEHTFEIKPLYEHLHAYVRAKLANAYPSYSP 240
Db 199 YGDYWRGDIYEEDTGYNSRSQLIKQVEHTFTQIKPLYQLHAYVRAKLANAYPSYSP 258
QY 241 IGCLPAHLGLDMWGRFTWNTLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSU 300
Db 259 TGCLPAHLGLDMWGRFTWNTLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSU 318
QY 301 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVWDDFLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNVOKAVCHPTAWDLGKGFRLMCTKVWDDFLTAHHEMGH 378
QY 361 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
Db 379 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438
QY 421 LLKQALTIIVGTLPTFTYMLEKRWVFKGEIPKQNMKKWEMKREIVGVVPEVPHDETYC 480
Db 439 LLKQALTIIVGTLPTFTYMLEKRWVFKGEIPKQNMKKWEMKREIVGVVPEVPHDETYC 498
QY 481 DPASLPHVSNDSYFRYYRTLYQFOFQALCOAAKHEGPLHKCDISNSTEAGOKLFNML 540
Db 499 DPASLPHVANDYSFIRYYRTIYQFOFQALCRKAKHEGPLHKCDISNSTEAGOKLFNML 558
QY 541 RLKSEPTWLTALENVVGAKNMVRPLNLYFPELFTWLKQDNKNSFVGWSTWDSY 595
Db 559 TLGKSKPWTLALEHVVGKKNVTPLLKYFPELFTWLKQDNKNSFVGWNTDWRPY 613

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RESULT 4
ACE2_PAGLA
ID ACE2_PAGLA STANDARD; PRT; 805 AA.
AC Q56NL1;
DT 13-SEP-2005 (Rel. 48, Created)
DT 13-SEP-2005 (Rel. 48, Last sequence update)
DT 13-SEP-2005 (Rel. 48, Last annotation update)
DE Angiotensin-converting enzyme 2 precursor (EC 3.4.17.-) (ACE-related
DE carboxypeptidase).
GN Name=ACE2;
OS Paguma larvata (Masked palm civet).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Laurasiatheria; Carnivora; Fissipedia; Viverridae;
OC Paradoxurinae; Paguma.
OX NCBI_TaxID=9675;
RN [1]
RP NUCLEOTIDE SEQUENCE, AND INTERACTION WITH HCOV-SARS S PROTEIN.
RX PubMed=15791205; DOI=10.1038/sj.emboj.7600640;
RA Li W., Zhang C., Sui J., Kunj J.H., Moore M.J., Luo S., Wong S.-K.,
RA Huang I.-C., Xu K., Vasiliava N., Murakami A., He Y., Marasco W.A.,
RA Guan Y., Choe H., Farzan M.;
RT "Receptor and viral determinants of SARS-coronavirus adaptation to
RT human ACE2.";
RL EMBO J. 24:1634-1643(2005).
CC -!- FUNCTION: Carboxypeptidase which converts angiotensin I to
CC angiotensin 1-9, a peptide of unknown function, and angiotensin II
CC to angiotensin 1-7, a vasodilator. Also able to hydrolyze apelin-
CC 13 and dynorphin-13 with high efficiency. May be an important
CC regulator of heart function (By similarity). Functional receptor
CC for human coronavirus SARS.
CC -!- COFACTOR: Binds 1 zinc ion per subunit (By similarity).
CC -!- COFACTOR: Binds 1 chloride ion per subunit (By similarity).
CC -!- SUBUNIT: Interacts with HCOV-SARS S protein.
CC -!- SUBCELLULAR LOCATION: Type I membrane protein. Processing by
CC ADAM17 may lead to a secreted protein (By similarity).
CC -!- SIMILARITY: Belongs to the peptidase M2 family.
CC
CC This Swiss-Prot entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use as long as its content is in no way modified and this statement is not
CC removed.
CC
CC EMBL; AY881174; AAX63775.1; -; mRNA.
CC InterPro; IPR010980; Cyt c b562.
CC InterPro; IPR006025; Pept M zn BS.
CC InterPro; IPR001548; Peptidase_M2.
CC Pfam; PF01401; Peptidase M2; 1.
CC PRINTS; PR00791; PEPTIDTASEA.
CC PROSITE; PS00142; ZINC_PROTEASE; 1.
CC Carboxypeptidase; Chloride; Glycoprotein; Hydrolase; Metal-binding;
CC Metalloprotease; Protease; Signal; Transmembrane; Zinc.
FT SIGNAL 1 17 Potential.
FT CHAIN 18 805 Angiotensin-converting enzyme 2.
FT TOPO_DOM 18 740 Extracellular (Potential).
FT TRANSMEM 741 761 Potential.
FT TOPO_DOM 762 805 Cytoplasmic (Potential).
FT REGION 30 41 Interaction with SARS S protein.
FT REGION 82 84 Interaction with SARS S protein (By
FT similarity).
FT REGION 90 93 Interaction with SARS S protein.
FT REGION 353 357 Interaction with SARS S protein (By
FT similarity).
FT ACT_SITE 375 375 By similarity.
FT ACT_SITE 505 505 Zinc (catalytic) (By similarity).
FT METAL 374 374 Zinc (catalytic) (By similarity).
FT METAL 378 378 Zinc (catalytic) (By similarity).
FT METAL 402 402 Zinc (catalytic) (By similarity).
FT BINDING 169 169 Chloride (By similarity).
FT BINDING 273 273 Substrate (By similarity).
FT BINDING 345 345 Substrate (By similarity).
FT BINDING 346 346 Substrate (By similarity).
FT BINDING 371 371 Substrate (By similarity).

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FT BINDING 477 477 Chloride (By similarity).
FT BINDING 481 481 Chloride (By similarity).
FT BINDING 515 515 Substrate (By similarity).
FT CARBOHYD 53 53 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 216 216 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 322 322 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 546 546 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 660 660 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 690 690 N-linked (GlcNAc...) (Potential).
FT DISULFID 133 141 By similarity.
FT DISULFID 344 361 By similarity.
FT DISULFID 530 542 By similarity.
SQ SEQUENCE 805 AA; 92611 MW; CF6406851F73E378 CRC64;

Query Match 86.5%; Score 2794; DB 1; Length 805;
Best Local Similarity 85.4%; Pred. No. 4.1e-189;
Matches 508; Conservative 36; Mismatches 51; Indels 0; Gaps 0;

QY 1 STIEQAKTFIDKFNHEADLFYQSSLASWYNTNTITENQNMNAGDKSAFLKEQST 60
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 19 STTEELAKTFLETENYEAQELSYQSSVASWYNTNTITDENAKNMNEAGAKWSAYYEBQSK 78
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 61 LAQMYPLQIEONLTVKLOLQALQQNGSSVLSGDSKRLNTILNTMTSTIYSTGKVCNPNP 120
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 79 LAQTYPLAEIQDAKIKRQLQALQQSGSSVLSADKSQRLNTILNMTSTIYSTGKVCNPNP 138
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 121 QECLLLEPGLNEMANSIDYNERLWAWESRSEVGKQLRPLYEYVVLKNEMARANHYED 180
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 139 QECLLLEPGLNIMENSKDYNERLWAWEGRAEYVGVKQLRPLYEYVVLKNEMARANNYED 198
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 181 YGDYWRGDEYGVNGVDYSGQLIEDYEHFEEIKPLYEHLHAYVRAKLMAYPSYISF 240
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 199 YGDYWRGDEYEEWTGGYNSRNQLIQDVEDTFEQIKPLYQHLHAYVRAKLMAYPSYISR 258
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 241 IGCLPAHLIGDMWGRFNTNLSLTVPGQKNIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 300
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 259 TGCLPAHLIGDMWGRFNTNLSLTVPGQKNIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 318
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAMDGLKGFRIILMCTKTWTDDFTAHHEMGH 360
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 319 GLPNMTQGFWNSMLTDPGDKGVKVCPTAMDGLKGFRIKMKCTKTWTDDFTAHHEMGH 378
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 361 IOYDMAYAAQFPLLRNGANEGFHEAVGIMSLSAATPHLKSIGLLSPDFQEDNTEINF 420
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 379 IOYDMAYAAQFPLLRNGANEGFHEAVGIMSLSAATPHLKTIGLLSPAFSDNTEINF 438
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 421 LLKQALTTVGTLPFTYMLEKRWVWYFKGSIKDOQWKWEMKREIVGVVEVPVHDETYC 480
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 439 LLKQALTTVGTLPFTYMLEKRWVWYFKGAIKPEQWQKRWEMKRNIVGVVEVPVHDETYC 498
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 481 DPASLFHVSNDYSFIRYTRTYLQFQOEALCQAAKHEGPHLKCDISNSTEAGOKLFNNML 540
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 499 DPASLFHVSNDYSFIRYTRTYLQFQOEALCQIAKHEGPHLKCDISNSTEAGOKLFNNML 558
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 541 RIGKSEPTLALENVVGAKNMNVRPLNYFPFLFTWLKDKQNKNSFVGWSTDSWSPY 595
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 559 SLGRSEPTLALERVVGAKNMNVRPLNYFPFLFTWLKDKQNKNSFVGWSTDSWSPY 613
DB ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 5
ACE2_RAT
ID ACE2_RAT STANDARD; PRT; 805 AA.
AC Q5EGZ1;
DT 13-SEP-2005 (Rel. 48, Created)
DT 13-SEP-2005 (Rel. 48, Last sequence update)
DT 13-SEP-2005 (Rel. 48, Last annotation update)
DE Angiotensin-converting enzyme 2 precursor (EC 3.4.17.-) (ACE-related
DE carboxypeptidase).
GN Name=Ace2;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi;
OC Muridae; Murinae; Rattus.

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QY 301 GLPNMTOGFENSLTDPGNVQKAVCHPTAWDILGKGFRLIMCTKYTMDDFLTAHHEMCH 360
 Db 319 GLPOMTPGFNTSMLEPFGDRKVVCHPTAWDILGKGFRLIMCTKYTMDDFLTAHHEMCH 378
 QY 361 IQYDMAAYAAQPFLLRNGANEGFHEAAGEIMSLSAATPKHLKSLGSLSPDQFQDNEITEINF 420
 Db 379 IQYDMAAYAAQPFLLRNGANEGFHEAAGEIMSLSAATPKHLKSLGSLSPDQFQDNEITEINF 438
 QY 421 LLKQALITIVGTLPTTYMLEKRWVWVFKGEIPKQDMKKMWMKREIVGVVPEVPHDETVC 480
 Db 439 LLKQALITIVGTLPTTYMLEKRWVWVFKGEIPKQDMKKMWMKREIVGVVPEVPHDETVC 498
 QY 481 DPASLFHVSNDYSFIRYRTYTLXQFOQALCOAAXHEGPHLHKCDISNSTEAGOKLFWML 540
 Db 499 DPASLFHVSNDYSFIRYRTYTLXQFOQALCOAAXHEGPHLHKCDISNSTEAGOKLFWML 558
 QY 541 RLKSGSPWTLALNENNVGAKNNVRPLNYPEPFTLWKDQNKNSFVGWSTDMSPY 595
 Db 559 SLGNSGSPWTLALNENNVGAKNNVRPLNYPEPFTLWKDQNKNSFVGWSTDMSPY 613
 RESULT 6
 ID ACE2 MOUSE STANDARD; PRT; 805 AA.
 AC QBR0T0; Q99N70; Q99N71;
 DT 13-SEP-2005 (Rel. 48, Created)
 DT 13-SEP-2005 (Rel. 48, Last sequence update)
 DE Angiotensin-converting enzyme 2 precursor (EC 3.4.17.-) (ACE-related carboxypeptidase).
 GN Name:ACE2;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi; Muroidea; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP NUCLEOTIDE SEQUENCE [MRNA] (ISOFORMS 1 AND 2), AND TISSUE SPECIFICITY. MEDLINE=22375506; PubMed=12487024; DOI=10.1080/1042517021000021608;
 RA Kurotsu T., Suzuki Y., Imai J., Sugano S., Hida M., Tanigami A., Muroi S., Yamada Y., Hanaoka K.;
 RT "Molecular cloning, mRNA expression, and chromosomal localization of mouse angiotensin-converting enzyme-related carboxypeptidase (mACE2).";
 RL DNA Seq. 13:217-220 (2002).
 RN [2]
 RP NUCLEOTIDE SEQUENCE [LARGE SCALE MRNA] (ISOFORM 1).
 RC STRAIN=FVB/N; TISSUE=Kidney;
 RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G., Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D., Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K., Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F., Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L., Stapleton M., Soares M.B., Donald M.F., Casavant T.L., Scheetz T.E., Brownstein M.J., Uedin T.B., Toshiyuki S., Carninci P., Prange C., Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J., Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H., Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W., Villalon D.K., Munz D.M., Sodergren E.J., Lu X., Gibbs R.A., Fahey J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A., Whiting M., Madan A., Young A.C., Shchepochenkov Y., Bouffard G.G., Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C., Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.N., Krzywinski M.I., Skalak U., Smailus D.E., Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 RN [3]
 RP FUNCTION.
 RX PubMed=12075344; DOI=10.1038/nature00786;
 RA Crackower M.A., Sarao R., Oudit G.Y., Yagil C., Kozieradzki I.,

RA Scanga S.E., Oliveira-dos-Santos A.J., da Costa J., Zhang L., Pei Y., Scholey J., Ferrario C.M., Manoukian A.S., Chappell M.C., Backx P.H., Yagil Y., Penninger J.M.;
 RT "Angiotensin-converting enzyme 2 is an essential regulator of heart function.";
 RL Nature 417:822-828 (2002).
 RN [4]
 RP FUNCTION.
 RX MEDLINE=22848473; PubMed=12967627; DOI=10.1016/S0022-2828(03)00177-9;
 RA Donoghue M., Wakimoto H., Maguire C.T., Acton S., Hales P., Stagliano N., Faichild-Huntress V., Xu J., Lorenz J.N., Kadambi V., Berul C.I., Breitbart R.E.;
 RT "Heart block, ventricular tachycardia, and sudden death in ACE2 transgenic mice with downregulated connexins.";
 RL J. Mol. Cell. Cardiol. 35:1043-1053 (2003).
 RN [5]
 RP INTERACTION WITH HCOV-SARS S PROTEIN.
 RX PubMed=15452268; DOI=10.1128/JVI.78.20.11429-11433.2004;
 RA Li W., Greenough T.C., Moore M.J., Vasileva N., Somasundaran M., Sullivan J.L., Farzan M., Choe H.;
 RT "Efficient replication of severe acute respiratory syndrome coronavirus in mouse cells is limited by murine angiotensin-converting enzyme 2.";
 RL J. Virol. 78:11429-11433 (2004).
 RN [6]
 RP TISSUE SPECIFICITY, INDUCTION, AND FUNCTION.
 RX PubMed=16001071; DOI=10.1038/nature03712;
 RA Imai Y., Kuba K., Rao S., Huan Y., Guo F., Guan B., Yang P., Sarao R., Wada T., Leong-Poi H., Crackower M.A., Fukamizu A., Hui C.-C., Hein L., Uhlig S., Slutsky A.S., Jiang C., Penninger J.M.;
 RT "Angiotensin-converting enzyme 2 protects from severe acute lung failure.";
 RL Nature 436:112-116 (2005).
 RN [7]
 RP TISSUE SPECIFICITY.
 RX PubMed=15949646; DOI=10.1016/j.peptides.2005.01.009;
 RA Gemardt F., Sterner-Kock A., Imboden H., Spalteholz M., Reibitz F., Orgathseiss H.-P., Siems W.-E., Walther T.;
 RT "Organ-specific distribution of ACE2 mRNA and correlating peptidase activity in rodents.";
 RL Peptides 26:1270-1277 (2005).
 CC -!- FUNCTION: Carboxypeptidase which converts angiotensin I to angiotensin 1-9, a peptide of unknown function, and angiotensin II to angiotensin 1-7, a vasodilator. Also able to hydrolyze apelin-13 and dynorphin-13 with high efficiency. May be an important regulator of heart function. May have a protective role in acute lung injury.
 CC -!- COFACTOR: Binds 1 zinc ion per subunit (By similarity).
 CC -!- COFACTOR: Binds 1 chloride ion per subunit (By similarity).
 CC -!- SUBUNIT: Weakly interacts with HCOV-SARS S protein.
 CC -!- SUBCELLULAR LOCATION: Type I membrane protein. Processing by ADAM17 may lead to a secreted protein (By similarity).
 CC -!- ALTERNATIVE PRODUCTS:
 CC Event-Alternative splicing; Named isoforms=2;
 CC Name=1;
 CC IsoId=QBR0T0-1; Sequences=Displayed;
 CC Name=2;
 CC IsoId=QBR0T0-2; Sequences=VSP 014903;
 CC -!- TISSUE SPECIFICITY: Expressed in heart, kidney and forebrain (at protein level). Ubiquitously expressed, with highest levels in ileum, kidney and lung. In lung, expressed on vascular endothelial and airway epithelial cells.
 CC -!- INDUCTION: Down-regulated in lung after acute injury.
 CC -!- MISCELLANEOUS: Mice lacking ACE2 are viable and fertile, exhibit normal kidney and lung function, but show a severe reduction in cardiac contractility, and are highly sensitive to severe acute lung failure. Transgenic mice overexpressing ACE2 in the heart appear healthy but show conduction disturbances and ventricular arrhythmias which can lead to sudden death.
 CC -!- CAUTION: Ref.1 (BAB40431) sequence differs from that shown due to a frameshift in position 784.

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DR InterPro: IPR001548; Peptidase_M2.
DR Pfam: PF01401; Peptidase_M2; 1.
DR PRINTS: PR00791; PEPDIPASEA.
DR PROSITE: PS00142; ZINC_PROTEASE; 1.
KW Carboxypeptidase; Chloride; Glycoprotein; Hydrolase; Metal-binding;
KW Metalloprotease; Protease; Signal; Transmembrane; Zinc.
FT SIGNAL 1 17
FT CHAIN 18 804
FT TOPO_DOM 18 739
FT TRANSMEM 740 760
FT TOPO_DOM 761 804
FT ACT_SITE 374 374
FT ACT_SITE 504 504
FT METAL 373 373
FT METAL 377 377
FT METAL 401 401
FT METAL 468 468
FT BINDING 168 168
FT BINDING 272 272
FT BINDING 344 344
FT BINDING 345 345
FT BINDING 370 370
FT BINDING 476 476
FT BINDING 480 480
FT BINDING 514 514
FT CARBOHYD 53 53
FT CARBOHYD 90 90
FT CARBOHYD 298 298
FT CARBOHYD 431 431
FT CARBOHYD 545 545
FT CARBOHYD 659 659
FT CARBOHYD 689 689
FT DISULFID 343 360
FT DISULFID 529 541
SQ SEQUENCE 804 AA; 93067 MW; E81570A96872A963 CRC64;

Query Match 83.9%; Score 2710.5; DB 1; Length 804;
Best Local Similarity 82.7%; Pred. No. 3.4e-183;
Matches 492; Conservative 48; Mismatches 54; Indels 1; Gaps 1;

QY 1 STIEQAKFLDKFNHEADLFQSSLASWNTNTITERNVQNMNAGDKWAFLEKQST 60
DB 19 STIEQAKFLDKFNHEADLFQSSLASWNTNTITERNVQNMNAGDKWAFLEKQST 78

QY 61 LAQMYPLOBIQNLTKVLQALQOQNGSSVLSDEKSKRLNTILNTMTSTYTGKVCNPNP 120
DB 79 MAKTSLEIQLNTLRQLKALQHSSTLSAESAESKRLNTILNTMTSTYTGKVLDP-NT 137

QY 121 QECLELLEPGLNIMANSLDYNRLWAWESWRSEVKGQLRPLRYEYVVLKQNMARANHED 180
DB 138 QECLELLEPGLDIDMENSRYNRLWAWEGWRAEVRGQLRPLRYEYVVLKQNMARANNED 197

QY 181 YGDYWRGDIYVNGVDGYDSRQLIEDVHTTEIEIKPLYLEHLHAYVRKLMNAYPSYISP 240
DB 198 YGDYWRGDIYVNGVDGYDSRQLIEDVHTTEIEIKPLYLEHLHAYVRKLMNAYPSYISP 257

QY 241 IGCPLPAHLGDMWGRFTWNLXSLTVPFGKPNIDVDAMVDQADWADQRIFFKAERFFVSV 300
DB 258 TGCPLPAHLGDMWGRFTWNLXSLTVPFEHKPSIDVTCKENQSWDAERIFKAERFFVSI 317

QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGDFFILMCTKVTMDDFLTAHMEGH 360
DB 318 SLPYMTQGFWDNSMLTEPGDGRKVVCHPTAWDLGKGDFFIKMCTKVTMDDFLTAHMEGH 377

QY 361 IQYDMAYAAQPELLRNGANEGHEAVGEIMSLSAATPKHLKSGILLSPDFQEDNETINF 420
DB 378 IQYDMAYAAQPYLLRNGANEGHEAVGEIMSLSAATPKHLKSGILLSPDFQEDNETINF 437

QY 421 LKQALTIYVGLTPTFTMLEKRWMMVFKGEIPKQNMKQWEMKREITGVVVEVPVPHDETVC 480
DB 438 LKQALTIYVGLTPTFTMLEKRWMMVFKGEIPKQNMKQWEMKREITGVVVEVPVPHDETVC 497

QY 481 DPASLPHVSNDSYFIRYTRTYIYQFQHEALCKTAKHEGALFKCDISNSTEAGQLQL 540
DB 481 DPASLPHVSNDSYFIRYTRTYIYQFQHEALCKTAKHEGALFKCDISNSTEAGQLQL 540

DB 498 DPACLFHVAEDYSFIRYTRTYIYQFQHEALCKTAKHEGALFKCDISNSTEAGQLQL 557
QY 541 RLIGKSEPTWTLALENVVVGAKNMNVRPLNFFFLFTLWKDQNKNSFVGVSTWSPY 595
DB 558 RLIGKSEPTWTLALENVVVGAKNMNVRPLNFFFLFTLWKDQNKNSFVGVSTWSPY 612

RESULT 8
ID Q5U380 BRARE PRELIMINARY; PRT; 785 AA.
AC Q5U380
DT 01-FEB-2005 (TRENBLrel. 29, Created)
DT 01-FEB-2005 (TRENBLrel. 29, Last sequence update)
DT 01-FEB-2005 (TRENBLrel. 29, Last annotation update)
DE Zgc:92514.
GN Name=ZGC:92514;
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP NUCLEOTIDE SEQUENCE.
RC TISSUE=Whole;
RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loguella N.A., Peters G.J., Alamek J.A., Gunaratne P.H.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.C., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettner M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalish D.E.,
RA Schnerch A., Schein J.E., Jones S.J.W., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
and mouse cDNA sequences";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
RN [2]
RP NUCLEOTIDE SEQUENCE.
RC TISSUE=Whole;
RG NIH MGC Project;
RL Submitted (NOV-2004) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC085667; AAB85667.1; -; mRNA.
DR ZFIN; ZDB-GENE-041114-6; ZGC:92514.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0004246; F:peptidyl-dipeptidase A activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR006025; Pept_M_Zn_BS.
DR InterPro; IPR001548; Peptidase_M2.
DR Pfam; PF01401; Peptidase_M2; 1.
DR PRINTS; PR00791; PEPDIPASEA.
DR ProDom; PD004184; Peptidase_M2; 1.
DR PROSITE; PS00142; ZINC_PROTEASE; UNKNOWN 1.
SQ SEQUENCE 785 AA; 90862 MW; 71CDF94B8772BDB1 CRC64;

Query Match 62.5%; Score 2018; DB 2; Length 785;
Best Local Similarity 61.4%; Pred. No. 4e-134;
Matches 366; Conservative 82; Mismatches 144; Indels 4; Gaps 3;

QY 2 TIEQAKFLDKFNHEADLFQSSLASWNTNTITERNVQNMNAGDKWAFLEKQSTL 61
DB 19 TVEDPAREFLNKFDEASDIMYQYTLASWAYNTDISQENADKEAYAINSYNKMSEE 78

QY 62 AQMYPLOBIQNLTKVLQALQOQNGSSVLSDEKSKRLNTILNTMTSTYTGKVCNPNP 121
DB 62 AQMYPLOBIQNLTKVLQALQOQNGSSVLSDEKSKRLNTILNTMTSTYTGKVCNPNP 121

Db 79 SNAYPIDQSDPIIKMQLQKQSGALSPDKASRLRNIMSEMSTIYNTATVCKIDDP 138
 QY 122 ECLLEPGELNETMANSLEDYERLWAWESWRSEVQKRLPLRYEYVVLKNEAMARANHYEDY 181
 Db 139 DCQTELEFGLSEMAESRDIDERLHWEGWRVATGMKORPLRYEYVVLKNEAKLNHYEDH 198
 QY 182 GDYWRGDYEVNGVDGYDGRGQIEDVEHTFEEIKPLYEHLHAYYRAKLMNAYPSYISPI 241
 Db 199 GDYWRGDYETIDDPKYSYSDQVIEDARRIYKEILLPLYKELHAYYRAKLMNAYPSYISPI 258
 QY 242 GCLPAHLGLDMGRFWTNLYSLTVPPGQKPNIDVTAMDQAWDAQRIFKEAEKPFVSVG 301
 Db 259 ACLPAHLGLDMGRFWTNLYSLTVPPGQKPNIDVTAMDQAWDAQRIFKEAEKPFVSVG 318
 QY 302 LNNMTQGFWENSLTDPGNVQXAVCHPTAWDLG-KGDFRILMCTKVWDDFLTAHEMCH 360
 Db 319 MPAMFDFNWNMSFIKP-BERDVVCHPTAWDMGNRDKFRIRKCTKVMDDFLTVHMHG 377
 QY 361 IOYDMAYAAQPFLLRNGANEGHEAVEIMSLSAATPKHLKSGILLSPDFQEDNTEINF 420
 Db 378 NQYQWAYRNHPYLLRNGANEGHEAVEIMSLSAATPSHLQSLGLPDKQDYETDINF 437
 QY 421 LKQALTIYVGLPFTYMLKRWMPFKGPIPKDQWKKWEMKREIVGVVEVPVPHDETYC 480
 Db 438 LKQALTIYVGLPFTYMLKRWMPFKGPIPKDQWKKWEMKREIVGVVEVPVPHDETYC 497
 QY 481 DPASLPHVSNDSYFIRYRTLYQFOQFALCOAAKHEGLHKCDISNSTEAGQKLFNWL 540
 Db 498 DPPALFHVSGDSYFIRYRTLYQFOQFALCOAAKHEGLHKCDISNSTEAGQKLFNWL 557
 QY 541 RLKKESEPTLALENVVGNKMMVRPLNLYFEPLFTWLKDQNK--NSFVGWSTDWSP 594
 Db 558 ELGRSNSTWALAEVAGTTKMSQPLHLYFSTLMEWLKEENKNNRVPCWVNVNP 613

RESULT 9
 Q4SHR0 TETNG
 ID Q4SHR0_TETNG PRELIMINARY; PRT; 652 AA.
 AC Q4SHR0;
 DT 13-SEP-2005 (TREMELrel. 31, Created)
 DT 13-SEP-2005 (TREMELrel. 31, Last sequence update)
 DT 13-SEP-2005 (TREMELrel. 31, Last annotation update)
 DE Chromosome 5 SCAFI4581, whole genome shotgun sequence.
 GN ORFNames=GSTENG0018041001;
 OS Tetraodon nigroviridis (Green puffer).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 OC Acanthomorpha; Acanthopterygii; Percormorpha; Tetraodontiformes;
 OC Tetraodontidae; Tetraodontidae; Tetraodon.
 OX NCBI_TaxID=99883;
 RN [1]
 RP NUCLEOTIDE SEQUENCE.
 RA Jallion O., Aury J.M., Brunet F., Petit J.L., Stange-Thomann N.,
 RA Mauceli E., Bouneau L., Fischer C., Ozouf-Costaz C., Bernot A.,
 RA Nicaud S., Jaffe D., Fisher S., Lutfalla G., Dossat C., Segurens B.,
 RA Dasilva C., Salanoubat M., Levy M., Boudet N., Castellano S.,
 RA Anthouard V., Jubin C., Castelli V., Katinka M., Vacherie B.,
 RA Blémond C., Skalli Z., Bottolico L., Poulain J., De Berardinis V.,
 RA Parra G., Lardier G., Chappie P., Coutanceau J.P., Gouzy J.,
 RA Kellis M., Volff J.N., Guigo R., Zody M.C., McKernan K.J., McEwan P., Bosak S.,
 RA Lindblad-Toh K., Birren B., Nusbaum C., Kahn D., Robinson-Rechavi M.,
 RA Laudet V., Schachner V., Quéfier F., Saurin W., Scarpelli C.,
 RA Wincker P., Lander E.S., Weissbach J., Roest Croallin H.;
 RT "Genome duplication in the teleost fish Tetraodon nigroviridis reveals
 the early vertebrate proto-karyotype.";
 RL Nature 431:946-957(2004).
 RN [2]
 RP NUCLEOTIDE SEQUENCE.
 RG Genoscope; Whitehead Institute Centre for Genome Research;
 RL Submitted (FEB-2004) to the EMBL/GenBank/DBJ databases.
 CC -!- CAUTION: The sequence shown here is derived from an
 EMBL/GenBank/DBJ whole genome shotgun (WGS) entry which is

CC preliminary data.
 DR EMBL; CAAB01014581; CAF99822.1; -, Genomic DNA.
 SQ SEQUENCE 652 AA; 75369 MW; 75784B3D18283309 CRC64;
 Query Match 49.9%; Score 1613; DB 2; Length 652;
 Best Local Similarity 64.2%; Pred. NO. 1.6e-105;
 Matches 299; Conservative 58; Mismatches 105; Indels 4; Gaps 3;
 QY 134 MANSIDYNERLWAWESWRSEVQKRLPLRYEYVVLKNEAMARANHYEDYDGYWRGDYEVNG 193
 Db 1 MANSIDYNERLWAWESWRSEVQKRLPLRYEYVVLKNEAMARANHYEDYDGYWRGDYEVNG 60
 QY 194 VD-GYDYSRGQIEDVEHTFEEIKPLYEHLHAYYRAKLMNAYPSYISPIGCLPAHLGLDM 252
 Db 61 EDPQPLYTRDELMDKDVRSAYKEILLPLYKELHAYYRAKLMNAYPSYISPIGCLPAHLGLDM 120
 QY 253 WGRFWTNLYSLTVPPGQKPNIDVTAMDQAWDAQRIFKEAEKPFVSVGLPMTQGFVEN 312
 Db 121 WGRFWTNLYSLTVPPGQKPNIDVTAMDQAWDAQRIFKEAEKPFVSVGLPMTQGFVEN 180
 QY 313 SMLTDPGNVQXAVCHPTAWDLG-KGDFRILMCTKVWDDFLTAHEMCHIOYDMAYAAQ 371
 Db 181 SMLTDPGNVQXAVCHPTAWDLG-KGDFRILMCTKVWDDFLTAHEMCHIOYDMAYAAQ 240
 QY 372 FLLRNGANEGHEAVEIMSLSAATPKHLKSGILLSPDFQEDNTEINFLLKQALTIYVGT 431
 Db 241 YPLRNGANEGHEAVEIMSLSAATPKHLKSGILLSPDFQEDNTEINFLLKQALTIYVGT 300
 QY 432 LPFTYMLKRWMPFKGPIPKDQWKKWEMKREIVGVVEVPVPHDETYCDPASLPHVSN 491
 Db 301 LPFTYMLKRWMPFKGPIPKDQWKKWEMKREIVGVVEVPVPHDETYCDPASLPHVSN 360
 QY 492 YSFIRYRTLYQFOQFALCOAAKHEGLHKCDISNSTEAGQKLFNMLRKGSPWTLA 551
 Db 361 YSFIRYRTLYQFOQFALCOAAKHEGLHKCDISNSTEAGQKLFNMLRKGSPWTLA 420
 QY 552 LENVVGNKMMVRPLNLYFEPLFTWL--KDQNKNSFVGWSTDWSPY 595
 Db 421 LKTSIGDVRMAAPLLDYFKLHDLWLVENKNNRNTVGMKTETEPY 466

RESULT 10
 ACE MESAU
 ID ACE MESAU STANDARD; PRT; 1314 AA.
 AC Q50JES;
 DT 13-SEP-2005 (Rel. 48, Created)
 DT 13-SEP-2005 (Rel. 48, Last sequence update)
 DT 13-SEP-2005 (Rel. 48, Last annotation update)
 DE Angiotensin-converting enzyme, somatic isoform precursor (EC 3.4.15.1)
 DE (Dipeptidyl carboxypeptidase I) (Kininase II) [Contains: Angiotensin-
 converting enzyme, somatic isoform, soluble form].
 GN Name=Ace; Synonyms=bcpl;
 OS Mesocricetus auratus (Golden hamster).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi;
 OC Muridae; Cricetidae; Cricetinae; Mesocricetus.
 OX NCBI_TaxID=10036;
 RN [1]
 RP NUCLEOTIDE SEQUENCE [MRNA].
 RA Uchida T.;
 RT "cDNA cloning of hamster angiotensin converting enzyme and mRNA
 expression in organs.";
 RL Submitted (MAY-2005) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: Converts angiotensin I to angiotensin II by release of
 the terminal His-Leu, this results in an increase of the
 vasoconstrictor activity of angiotensin. Also able to inactivate
 bradykinin, a potent vasodilator (By similarity).
 CC -!- CATALYTIC ACTIVITY: Release of a C-terminal dipeptide,
 oligopeptide--Xaa-Yaa, when Xaa is not Pro, and Yaa is neither
 Asp nor Glu. Thus, conversion of angiotensin I to angiotensin II,
 with increase in vasoconstrictor activity, but no action on
 angiotensin II.
 CC -!- COPACTOR: Binds 2 zinc ions per subunit (By similarity).
 CC


```
FT DISULPID 569 581 S -> P (in dbSNP:4317).
FT VARIANT 32 32 /FTId=VAR 011710.
FT 49 S -> G (in dbSNP:4318).
FT VARIANT 49 49 /FTId=VAR 011711.
FT 342 T -> M (in dbSNP:3730043).
FT VARIANT 342 342 /FTId=VAR 023435.
FT 444 I -> T (in dbSNP:4976).
FT VARIANT 444 444 /FTId=VAR 014738.
FT 477 F -> V (in dbSNP:4977).
FT VARIANT 477 477

Query Match 41.3%; Score 1335; DB 1; Length 732;
Best Local Similarity 41.9%; Pred. No. 9.4e-86;
Matches 254; Conservative 116; Mismatches 202; Indels 34; Gaps 9;

QY 2 TIEBAKTFELDKFNHEADLFYQSSLSASWNTNTITEE-----NVQNMNAGDKWSA 53
Db 70 TDEABASKFVEYDRTSQVWVNEAENWNTNTITTSKILLQKNQIANHT----- 123
QY 54 FLKEQSTLAQMPLOEIQNLTVKLQALQOQNGSSVLSEDKSKRLNTILNTMTSTIYSTGK 113
Db 124 --LKYGTOARKFDVNLQNTTKRIKKVQDLERAAQAELEEFNKILLDMETTSVAT 181
QY 114 VCNPDNPOECLELLEPGLNEIMANSIDYNERLWAMESRSEVGKQLRPLYEYVVLKNEMA 173
Db 182 VCHPNG--SCLEPEDLTNWMATSKYEDLLWAMEGRDKAGRAILQFPKYVELINQAA 239
QY 174 RANHVEDYGDYWRGDIYVNGVDYDYSRGQIEDVEHTPEEIKPLYEHLHAYVRAKLMA 233
Db 240 RLNGYVDAGDSWRSWYETPSLE-----QDLERLFOELQPLLYNLHAYVRRALHRH 289
QY 234 Y-PSYISPIGCLPAHLLGDMGRFWNTLYSLTVPGQKPNIDVTAMVDQAWDAQRIKFE 292
Db 290 YGAQHINLEGPPIPAHLLGNMWAQTSNIYDLVVPFAPSMDTTEAMLKQGWTPRRMFKE 349
QY 293 AEKPFVSVGLPNMTOGFWNSMLTDPGNVQKAVCHPTAWDLGK--DFRILMCTKVTMDDF 351
Db 350 ADDFTSLGLLPVPEFWNKSMLERKPTDGREVVVCHASAWDFYNGKDFRIKQCTTVNLEDL 409
QY 352 LTAHEMHGHIQDYMAAOPFLLRGANEGFHEAAGEIMSLSAATPKHLKSLGLLSPDFQ 411
Db 410 VVAHEMHGHIQFYMQYKDLPLVALREGANPGFHEAGDVLALSVPKHLKSLNLSSEGG 469
QY 412 ENETETINFLKQALITVGLTPPTMLEKRWVFKGEIPKQWKKWEMKREIVGVVE 471
Db 470 SD-EHDINFLMKWALDKIAFIPFSLVDQWRVFDGSITKENYQWMSLRKYLQGLCP 528
QY 472 PVPHEITYCDPASLHVSNDSYFIRYVTLTYQFOFQALCOAAXHEGLHKCDISNTE 531
Db 529 PVPRTQGDPDFGAKFHPISVPIRYFVFIIFQFQFHEALCQAAGHTGPLHKCDIYQSKE 588
QY 532 AGQKLFNMLRLGKSEPTLALENVGAKNMVRLINYPEPLFTWLKQNK--NSFVGVW- 588
Db 589 AGQRLATAMKLGFSRWPPEAMQLITQPNWSASAMLSYKPLLDMLRTENELHGEKLGWP 648
QY 589 STDWSP 594
Db 649 QVNWTP 654

RESULT 12
QBN710 HUMAN PRELIMINARY; PRT; 739 AA.
AC QBN710.
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Angiotensin I converting enzyme, isoform 3.
GN Name=ACE;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae;
OC Homo.
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OX NCBI_TaxID=9606;
RN [1]
RP NUCLEOTIDE SEQUENCE.
RX MEDLINE=23388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
TISSUE-Testis;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
Klausner R.D., Collins P.S., Wagner L., Shenmen C.M., Schuler G.D.,
Altechul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
Stapleton M.J., Udén T.B., Toshiyuki S., Carninci P., Prange C.,
Raha S.S., Lequellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
Fahy J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smallus D.E.,
Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
"Generation and initial analysis of more than 15,000 full-length human
and mouse cDNA sequences.";
Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP NUCLEOTIDE SEQUENCE.
RC TISSUE-Testis;
RA Director MGC Project;
RL Submitted (AUG-2002) to the EMBL/GenBank/DBSJ databases.
DR EMBL; BC036375; AAH36375.1; -, mRNA.
DR HSSP; Q10714; 1J37.
DR SMR; QBN710; 71-648.
DR Ensembl; ENSG00000159640; Homo sapiens.
DR GO; GO:0016020; C:membrane; IEA.
DR GO; GO:0004246; F:peptidyl-dipeptidase A activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR001548; Peptidase M2.
DR Pfam; PF01401; Peptidase M2; 1.
DR PRINTS; PR00791; PEPTIDPASEA.
DR ProDom; PD004184; Peptidase M2; 1.
DR PROSITE; PS00142; ZINC PROTEASE; UNKNOWN 1.
SQ SEQUENCE 739 AA; 83958 MW; 87995DFP58D93D01 CRC64;
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Query Match 41.3%; Score 1335; DB 2; Length 739;
Best Local Similarity 41.9%; Pred. No. 9.5e-86;
Matches 254; Conservative 116; Mismatches 202; Indels 34; Gaps 9;
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QY 2 TIEBAKTFELDKFNHEADLFYQSSLSASWNTNTITEE-----NVQNMNAGDKWSA 53
Db 70 TDEABASKFVEYDRTSQVWVNEAENWNTNTITTSKILLQKNQIANHT----- 123
QY 54 FLKEQSTLAQMPLOEIQNLTVKLQALQOQNGSSVLSEDKSKRLNTILNTMTSTIYSTGK 113
Db 124 --LKYGTOARKFDVNLQNTTKRIKKVQDLERAAQAELEEFNKILLDMETTSVAT 181
QY 114 VCNPDNPOECLELLEPGLNEIMANSIDYNERLWAMESRSEVGKQLRPLYEYVVLKNEMA 173
Db 182 VCHPNG--SCLEPEDLTNWMATSKYEDLLWAMEGRDKAGRAILQFPKYVELINQAA 239
QY 174 RANHVEDYGDYWRGDIYVNGVDYDYSRGQIEDVEHTPEEIKPLYEHLHAYVRAKLMA 233
Db 240 RLNGYVDAGDSWRSWYETPSLE-----QDLERLFOELQPLLYNLHAYVRRALHRH 289
QY 234 Y-PSYISPIGCLPAHLLGDMGRFWNTLYSLTVPGQKPNIDVTAMVDQAWDAQRIKFE 292
Db 290 YGAQHINLEGPPIPAHLLGNMWAQTSNIYDLVVPFAPSMDTTEAMLKQGWTPRRMFKE 349
QY 293 AEKPFVSVGLPNMTOGFWNSMLTDPGNVQKAVCHPTAWDLGK--DFRILMCTKVTMDDF 351
Db 350 ADDFTSLGLLPVPEFWNKSMLERKPTDGREVVVCHASAWDFYNGKDFRIKQCTTVNLEDL 409
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RA Johnston C.I.;
 RT "Myocardial infarction increases ACE2 expression in rat and humans.";
 RL Eur. Heart J. 26:369-375(2005).
 RN [15]
 RP VARIANTS THR-1018; VAL-1051; GLN-1279; SER-1286 AND PRO-1296.
 RX MEDLINE=99318094; PubMed=10391210; DOI=10.1038/10297;
 RA Halushka M.K., Fan J.-B., Bentley K., Hsie L., Shen N., Weder A.,
 Cooper R., Lipshutz R., Chakravarti A.;
 RT "Patterns of single-nucleotide polymorphisms in candidate genes for
 blood-pressure homeostasis";
 RL Nat. Genet. 22:239-247(1999).
 CC -I- FUNCTION: Converts angiotensin I to angiotensin II by release of
 the terminal His-Leu; this results in an increase of the
 vasoconstrictor activity of angiotensin. Also able to inactivate
 bradykinin, a potent vasodilator.
 CC -I- CATALYTIC ACTIVITY: Release of a C-terminal dipeptide,
 oligopeptide-[Xaa-Yaa, when Xaa is not Pro, and Yaa is neither
 Asp nor Glu. Thus, conversion of angiotensin I to angiotensin II,
 with increase in vasoconstrictor activity, but no action on
 angiotensin II.
 CC -I- COFACTOR: Binds 2 zinc ions per subunit (By similarity).
 CC -I- COFACTOR: Binds 2 chloride ions per subunit (By similarity).
 CC -I- ENZYME REGULATION: Strongly activated by chloride. Specifically
 inhibited by lisinopril, captopril and enalaprilat.
 CC -I- BIOPHYSICOCHEMICAL PROPERTIES:
 CC Kinetic parameters:
 CC KM=2.51 mM for Hip-His-Leu;
 CC -I- SUBCELLULAR LOCATION: Type I membrane protein. A soluble form
 released by proteolysis also exists.
 CC -I- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=Somatic;
 CC IsoId=P12821-1; Sequence=Displayed;
 CC Name=Testis-specific;
 CC IsoId=P22966-1; Sequence=External;
 CC -I- TISSUE SPECIFICITY: Ubiquitously expressed, with highest levels in
 lung, kidney, heart, gastrointestinal system and prostate.
 CC -I- INDUCTION: Up-regulated in failing heart.
 CC -I- PTM: Phosphorylated by CK2 on Ser-1299, which allows membrane
 retention.
 CC -I- MISCELLANEOUS: Inhibitors of ACE are commonly used to treat
 hypertension and some types of renal and cardiac dysfunction.
 CC -I- SIMILARITY: Belongs to the peptidase M2 family.
 CC This Swiss-Prot entry is copyright. It is produced through a collaboration
 between the Swiss Institute of Bioinformatics and the EMBL outstation -
 the European Bioinformatics Institute. There are no restrictions on its
 use as long as its content is in no way modified and this statement is not
 removed.
 CC
 CC EMBL; J04144; AA51684.1; -; mRNA.
 CC EMBL; AF118569; AAD28560.1; -; Genomic DNA.
 CC EMBL; AY436326; AAR03504.1; -; Genomic DNA.
 CC EMBL; AB208971; BAD92208.1; ALT_INIT; mRNA.
 CC FIR; A31759; A31759.
 CC HSP; Q10714; IJ36.
 CC SNR; P12821; 645-1222.
 CC MEROPS; M02.001; -.
 CC MEROPS; M02.004; -.
 CC Ensembl; ENSG00000159640; Homo sapiens.
 CC HGNC; HGNC:2707; ACE.
 CC MIM; 106180; -.
 CC GO; GO:0005624; C:membrane fraction; TAS.
 CC GO; GO:0005886; C:plasma membrane; TAS.
 CC GO; GO:0005625; C:soluble fraction; TAS.
 CC GO; GO:0008217; P:regulation of blood pressure; TAS.
 CC InterPro; IPR006025; Pept M.2n.BS.
 CC Pfam; PF01401; Peptidase M2; 2.
 CC PRINTS; PR00791; PEPDPTASEA.
 CC ProDom; PD004184; Peptidase M2; 2.
 CC PROSITE; PS00142; ZINC PROTEASE; 2.
 CC Alternative splicing; Carboxypeptidase; Direct protein sequencing;
 KW

KW Glycoprotein; Hydrolase; Metal-binding; Metalloprotease;
 KW Phosphorylation; Polymorphism; Protease; Repeat; Signal;
 KW Transmembrane; Zinc.
 FT SIGNAL 1 29
 Query Match 41.3%; Score 1335; DB 1; Length 1306;
 Best Local Similarity 41.9%; Pred. No. 2.1e-85;
 Matches 254; Conservative 116; Mismatches 202; Indels 34; Gaps 9;
 QY 2 TTEQAKTFLDKFNHEADLFYQSSLASNNYNTTEE-----NVQNNNAGDKWSA 53
 Db TDEASAKFVEEDRTSQVWNEYAEANNVYNTTITSTSKILLQNMQIANHT----- 697
 QY 54 FLKEOSTLAQMPLOEIQNLTKVLQALQONGSSVLSEDKSKRLNTILNTSTIYSTGK 113
 Db 698 --LKYGTQARKFDVNLQNTTKRIKKVQDLERAAALPAQELBEYNKILLDMETTSVAT 755
 QY 114 VCNPNPQECILLLEPLNEIMANSIDYNERLWAWESRSEVGKQLRPLEYEYVVLKNEMA 173
 Db 756 VCHPNG--SCQLQEPDLTNWATSKRYEDLLWAWESGRDKAGRAILOQFPKYVELINQAA 813
 QY 174 RANHYEDYGYWRGDFYVNGVDGYDSRGQLIEDVEHTFEETKPLYEHLHAYVRAKLMA 233
 Db 814 RLNGYVDAGDSWRSWYETPSLE-----QDLERLFQELQPLLYNLHAYVRRALHRH 863
 QY 234 Y-PSYISPIGCLPAHLGDMGRFWTNLYSLTPFGQKFNIDVTDMVDQAWDAQRIKFE 292
 Db 864 YGAQHINLEGPITPAHLGLGNMAQTWSNIYDLVVPFSPASMDTTEAMLKQGTPTRRMFE 923
 QY 293 AEKFFVSGLPNMTQGFWENSMITDPGNVOKAVCHPTAWDLGKG-DFTLMCTKVTMDDF 351
 Db 924 ADDFTSLGLLPVPPEFWNKSMLEKPTDQREVVCASAWDFYNGKDFRIKQCTTVNLEDL 983
 QY 352 LTAHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSLGLSPDFQ 411
 Db 984 VVAHEMGHIQYFMQYKDLPAVALREGANPGFHEAIGDVLALSVSTPKHLSLNLSSEGG 1043
 QY 412 ENETETINFLKQALTYIGTLPFTTMLEKRWMMFKGEIPKQWMMKWMRELVGVVE 471
 Db 1044 SD-EHDINFLMKVADLKIAFTPSYLVQMRWRVFDGSIITKENYQWMSLRLKYGGLCP 1102
 QY 472 VPVHDETCDPASLPHVSNDSYFIRYTRTLVQFOFQALCOAAKHGPHLKCDSNSTE 531
 Db 1103 PVPRTQGFDPGAKPHIPISSVPYIRYFVSFIQFQHEALCOAAGHTGPLHKCDIYQSK 1162
 QY 532 AQQLFNMLRLGKSPFTLALENVVYAKMNVRLPLNTFPEPLFTWLKDQNK--NSFVGVW- 588
 Db 1163 AQQLATAMKLGFSRPWEPMQLITQPNWSASAMLSYFKPLLDMLRTENELHGEKLGWP 1222
 QY 589 STDWSP 594
 Db 1223 QYNWTP 1228
 RESULT 14
 ACET MOUSE
 ID ACET MOUSE STANDARD; PRT; 732 AA.
 AC P22967;
 DT 01-AUG-1991 (Rel. 19, Created)
 DT 01-AUG-1991 (Rel. 19, Last sequence update)
 DT 13-SEP-2005 (Rel. 48, Last annotation update)
 DE Angiotensin-converting enzyme, testis-specific isoform precursor
 DE (EC 3.4.15.1) (EC 3.2.1.-) (ACE-T) (Dipeptidyl carboxypeptidase I)
 DE (Kinase II) [Contains: Angiotensin-converting enzyme, testis-
 specific isoform, soluble form].
 GN Name=Ace; Synonyms=Dcpl;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi;
 OC Muridea; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP NUCLEOTIDE SEQUENCE [GENOMIC DNA / MRNA].

RX MEDLINE=90318396; PubMed=2164636;
 RA Howard T.E., Shai S.-Y., Langford K.G., Martin B.M., Bernstain K.E.;
 RT "Transcription of testicular angiotensin-converting enzyme (ACE) is
 RL initiated within the 12th intron of the somatic ACE gene."; Mol. Cell. Biol. 10:4294-4302(1990).
 RN [2]
 RX FUNCTION.
 RP PubMed=7753170; DOI=10.1038/375146a0;
 RA Krege J.H., John S.W., Langenbach L.L., Hodgkin J.B., Hageman J.R.,
 RA Bachman E.S., Jennette J.C., O'Brien D.A., Smithies O.;
 RT "Male-female differences in fertility and blood pressure in ACE-
 RL deficient mice."; Nature 375:146-148(1995).
 RN [3]
 RX FUNCTION.
 RP PubMed=8642790; DOI=10.1038/nm1179;
 RA Esther C.R. Jr., Howard T.E., Marino E.M., Goddard J.M.,
 RA Capocchi M.R., Bernstein K.E.;
 RT "Mice lacking angiotensin-converting enzyme have low blood pressure,
 RL renal pathology, and reduced male fertility."; Lab. Invest. 74:953-965(1996).
 RN [4]
 RX FUNCTION, ENZYME REGULATION, AND MUTAGENESIS OF HIS-413; GLU-414 AND
 RP HIS-417.
 RP PubMed=15665832; DOI=10.1038/nm1179;
 RA Kondoh G., Tojo H., Nakatani Y., Komazawa N., Murata C., Yamagata K.,
 RA Maeda Y., Kinoshita T., Okabe M., Taguchi R., Takeda J.;
 RT "Angiotensin-converting enzyme is a GPI-anchored protein releasing
 RL factor crucial for fertilization."; Nat. Med. 11:160-166(2005).
 CC -!- FUNCTION: Converts angiotensin I to angiotensin II by release of
 CC the terminal His-Leu, this results in an increase of the
 CC vasoconstrictor activity of angiotensin. Also able to inactivate
 CC bradykinin, a potent vasodilator. Has also a glycosidase activity
 CC which releases GPI-anchored proteins from the membrane by cleaving
 CC the mannose linkage in the GPI moiety. This GPIase activity seems
 CC to be crucial for the egg-binding ability of the sperm.
 CC -!- CATALYTIC ACTIVITY: Release of a C-terminal dipeptide,
 CC oligopeptide-I-Xaa-Yaa, when Xaa is not Pro, and Yaa is neither
 CC Asp nor Glu. Thus, conversion of angiotensin I to angiotensin II,
 CC with increase in vasoconstrictor activity, but no action on
 CC angiotensin II.
 CC -!- COFACTOR: Binds 2 zinc ion per subunit (By similarity).
 CC -!- COFACTOR: Binds 2 chloride ions per subunit (By similarity).
 CC -!- ENZYME REGULATION: Peptidase activity is specifically inhibited by
 CC lisinopril, captopril and enalaprilat. In contrast, GPIase
 CC activity is nearly insensitive to captopril.
 CC -!- SUBCELLULAR LOCATION: Type I membrane protein. A soluble form
 CC released by proteolysis also exists (By similarity).
 CC -!- ALTERNATIVE PRODUCTS.
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=Testis-specific;
 CC IsoId=P22967-1; Sequences=Displayed;
 CC Name=Somatic;
 CC IsoId=P09470-1; Sequences=External;
 CC -!- TISSUE SPECIFICITY: Spermatoocytes, adult testis.
 CC -!- INDUCTION: Expression is thought to be subject to hormonal
 CC regulation by androgens.
 CC -!- PTM: Phosphorylated by CK2 on Ser-725; which allows membrane
 CC retention (By similarity).
 CC -!- MISCELLANEOUS: Mice lacking ACE have low blood pressure, elevated
 CC serum potassium, anemia, and renal defects. Male mice have reduced
 CC fertility.
 CC -!- SIMILARITY: Belongs to the peptidase M2 family.
 CC -----
 CC This Swiss-Prot entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use as long as its content is in no way modified and this statement is not
 CC removed.
 CC -----
 CC EMBL; M55333; AAA37149.1; -; mRNA.
 CC EMBL; M61094; AAA37150.1; -; Genomic_DNA.

DR PIR; A35655; A35655.
 DR HSSP; Q10714; IJ36.
 DR SMR; P22967; 70-647.
 DR Ensembl; ENSMUSG0000020681; Mus musculus.
 DR MGI; MGI:87874; Ace.
 DR GO; GO:0005615; C:extracellular space; TAS.
 DR GO; GO:0016021; C:integral to membrane; TAS.
 DR InterPro; IPR006025; Pept_M_Zn_BS.
 DR InterPro; IPR001548; Peptidase_M2.
 DR Pfam; PF01401; Peptidase_M2_1.
 DR PRINTS; PR00791; PEPDIPASEA.
 DR ProDom; PD004184; Peptidase_M2; 1.
 DR PROSITE; PS00142; ZINC_PROTEASE; 1.
 KW Glycosidase; Hydrolase; Metal-binding; Metalloprotease;
 KW Phosphorylation; Protease; Signal; Testis; Transmembrane; Zinc.
 FT SIGNAL 1 31 By similarity.
 FT CHAIN 32 732 Angiotensin-converting enzyme, testis-specific isoform.
 FT FT Angiotensin-converting enzyme, testis-specific isoform.
 FT FT Removed in secreted form (By similarity).
 FT FT Extracellular (Potential).
 FT FT Cytoplasmic (Potential).
 FT FT Potential.
 FT FT Zinc (catalytic) (By similarity).
 FT FT Zinc (catalytic) (By similarity).
 FT FT Zinc (catalytic) (By similarity).
 FT FT Chloride 1 (By similarity).
 FT FT Chloride 2 (By similarity).
 FT FT Chloride 1 (By similarity).
 FT FT Chloride 1 (By similarity).
 FT FT Chloride 2 (By similarity).
 FT FT Phosphoserine (By similarity).
 FT FT N-linked (GlcNAc...) (Potential).
 FT FT N-linked (GlcNAc...) (complex) (By similarity).
 FT FT N-linked (GlcNAc...) (complex) (By similarity).
 FT FT N-linked (GlcNAc...) (complex) (By similarity).
 FT FT N-linked (GlcNAc...) (Potential).
 FT FT N-linked (GlcNAc...) (Potential).
 FT FT By similarity.
 FT FT By similarity.
 FT FT By similarity.
 FT FT H->K; Abolishes peptidase activity but no effect on GPIase activity; when associated to K-417.
 FT FT E->D; Abolishes peptidase activity but no effect on GPIase activity.
 FT FT H->K; Abolishes peptidase activity but no effect on GPIase activity; when associated to K-413.
 FT FT SEQUENCE 732 AA; 84047 MW; 16C817E7FBD09BD9 CRC64;
 Query Match 41.3%; Score 1334; DB 1; Length 732;
 Best Local Similarity 42.6%; Pred. No. 1.1e-85;
 Matches 255; Conservative 112; Mismatches 213; Indels 18; Gaps 7;
 QY 2 TIEQAKTFLDKNHEAEDLFVQSSLASWYNTNTEENVNMMNAGKWSAFLKEQSTL 61
 Db 69 TDEAKADRFVEYDRTAQVLLNEVYAEANQVNTNITIEGSKILLEKSEVSNHTLK YGR 128
 QY 62 AQMPYEQIQLTKVLQALQONGSVLSKSKRLNTILNTMTSTIYTGKVCNPDNPQ 121
 Db 129 AKTFDVSFNQSSIKRIKKQLNLDRAVLPPKELBEYNQILLDMETVYSLNSCYTNG-- 186
 QY 122 ECLLEPGELNIMNSLDYNERLWAWESWRSEVQKPLPYEEYVVLKSNARAHYEDY 181
 Db 187 TCMLEPDLTNMATSRYKEILLWAKSWRKDKVGRAILFPFPFKYEFESNKAIAKNGYTD 246


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QY 182 GDYWRGDYVNGVDYDSRGQLIEDVHTFBEIKPLYEHLHAYVRKLMNAYPS-Y1SP 240
Db 247 GDSWRSLYESDNLSE-----QDLKLYQLQLYLNHLHAYVRRSLHRYGSEYINL 296
QY 241 IGLPAHLGLDMGRFWNTLYSLVTFGQKPNIDVTDAWQDAQDAQRIKFAEAEKFFVSV 300
Db 297 DGPPIAHLGLNWAQTSNIYDLVAPFPAPNIDATEAMIKQGTWPRRIFKEADNFTSL 356
QY 301 GLPNMTQGFWSNMLTDPGNQKAVCHPTANDLGKG-DPRILMCTKVWMDFLTAHHMG 359
Db 357 GLLPVPEPFNWSKLEKPTDGRVVCVCHPSAMDFTYNGKDFRIKQCTSVNMEDLVIAHHMG 416
QY 360 HLOYDMAYAAQPLLRNGANEGFHEAVGRIMSLSAATPKHLKSLGILSPDFQEDNETEIN 419
Db 417 HIQYFMQYKDLPTVTFREGANPGFHEAIGDIMALSSTPKHLYSLLNLSLE-MSGVEYDIN 475
QY 420 FLKQALITVGLTPFTYMLEKRWMMVFKGEIPKQDMKKWMMKREIVGVVFPVPHDET 479
Db 476 FLMKALDKIAFIPFSYLDQWRVDFDSITKENYQEWMSLRKYQGLCPVPRSQGD 535
QY 480 CDPASLFHVSNDYSFIRYTRTYQFOEALCOAAKHEGPHLKCDISNSTAGQKLFNM 539
Db 536 FDPGSKFHPVNPVYRVFSPFIQFQFHEALCRAAGHTGPLHKCDIYQSKAGKLLADA 595
QY 540 LRLGKSEPTWLALENVVGAKNNVRPLNYPFLPTWLDQNK--NSFVGM-STDWSP 594
Db 596 MKLGYSKPWEPEAMKLTGQPNMSASAMNYPFKPLTEWLTENRRHGETLGNPEYNWAP 653

RESULT 15
Q8K233 MOUSE
ID Q8K233_MOUSE PRELIMINARY; PRT; 1015 AA.
AC Q8K233;
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Ace protein (Fragment).
GN Name=Ace;
OS Mus musculus (Mouse).
OC Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi;
OC Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP NUCLEOTIDE SEQUENCE.
RC STRAIN=FVB/N; TISSUE=Mammary tumor. C3;
RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh P.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Uedin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
RA Bosak S.A., McSwain P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Ketterman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalios D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
RN [2]
RP NUCLEOTIDE SEQUENCE.
RC STRAIN=FVB/N; TISSUE=Mammary tumor. C3;
RA Strausberg R.;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC034367; AAH34367.1; -; mRNA.

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DR HSP; Q10714; L337.
DR SMR; Q8K233; 416-993.
DR MGI; MGI:87874; Ace.
DR GO; GO:0005615; C:extracellular space; TAS.
DR GO; GO:0016021; C:integral to membrane; TAS.
DR InterPro; IPR001548; Peptidase M2.
DR InterPro; IPR006025; Pept_M_Zn_BS.
DR Pfam; PF01401; Peptidase_M2_2.
DR PRINTS; PR00791; PEPDIPTASEA.
DR ProDom; PD004184; Peptidase M2; 2.
DR PROSITE; PS00142; ZINC_PROTEASE; UNKNOWN_2.
FT NON_TER 1
SQ SEQUENCE 1015 AA; 117509 MM; D097F69585553C27 CRC64;

Query Match 41.3%; Score 1334; DB 2; Length 1015;
Best Local Similarity 42.6%; Pred. No. 1.7e-85;
Matches 255; Conservative 112; Mismatches 213; Indels 18; Gaps 7;

QY 2 TIEEAKTFLOKFNHEADLFYQSSLASNYNTNITEENVQNMNAGDKWSAFLKEQSTL 61
Db 415 TDEAKADRFVEEYDRTAQVLLNEYAEANWQYNTNITIEGSKILLEKSTEVSHHTUKYGR 474
QY 62 AQMYPLQBIQNLTVKLQALQONGSSVLSEDKSLRNLNTILNTMTSTIYSTGKVCNPDNPQ 121
Db 475 AKTFVDSNFQSSIKRIIKQLNLDRAVLPPKELEYNQILLDMETTYLSNICYNG-- 532
QY 122 ECLLEPEGLNEIMANSLDYNERLMAWESWRSEVGKQLRPLYEEYVVLKVMEMAPANHEDY 181
Db 533 TCMPLEPLDTNNMATSRYKEELLMAKSWRDKVGRAILPFPKPYVEFSNKIAKNGYTDA 592
QY 182 GDYWRGDYVNGVDYDSRGQLIEDVHTFBEIKPLYEHLHAYVRKLMNAYPS-Y1SP 240
Db 593 GDSWRSLYESDNLSE-----QDLKLYQLQLYLNHLHAYVRRSLHRYGSEYINL 642
QY 241 IGLPAHLGLDMGRFWNTLYSLVTFGQKPNIDVTDAWQDAQDAQRIKFAEAEKFFVSV 300
Db 643 DGPPIAHLGLNWAQTSNIYDLVAPFPAPNIDATEAMIKQGTWPRRIFKEADNFTSL 702
QY 301 GLPNMTQGFWSNMLTDPGNQKAVCHPTANDLGKG-DPRILMCTKVWMDFLTAHHMG 359
Db 703 GLLPVPEPFNWSKLEKPTDGRVVCVCHPSAMDFTYNGKDFRIKQCTSVNMEDLVIAHHMG 762
QY 360 HLOYDMAYAAQPLLRNGANEGFHEAVGRIMSLSAATPKHLKSLGILSPDFQEDNETEIN 419
Db 763 HIQYFMQYKDLPTVTFREGANPGFHEAIGDIMALSSTPKHLYSLLNLSLE-MSGVEYDIN 821
QY 420 FLKQALITVGLTPFTYMLEKRWMMVFKGEIPKQDMKKWMMKREIVGVVFPVPHDET 479
Db 822 FLMKALDKIAFIPFSYLDQWRVDFDSITKENYQEWMSLRKYQGLCPVPRSQGD 881
QY 480 CDPASLFHVSNDYSFIRYTRTYQFOEALCOAAKHEGPHLKCDISNSTAGQKLFNM 539
Db 882 FDPGSKFHPVNPVYRVFSPFIQFQFHEALCRAAGHTGPLHKCDIYQSKAGKLLADA 941
QY 540 LRLGKSEPTWLALENVVGAKNNVRPLNYPFLPTWLDQNK--NSFVGM-STDWSP 594
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Search completed: March 28, 2006, 11:16:07
Job time : 126.785 secs

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GenCore version 5.1.7
Copyright (c) 1993 - 2006 Bioacceleration Ltd.

OM protein - protein search, using sw model

Run on: March 28, 2006, 11:36:19 ; Search time 95.1865 Seconds
(without alignments)
2611.802 Million cell updates/sec

Title: US-10-659-000-4
Perfect score: 3231
Sequence: 1 STIEEQAKTFLDKFNHEAD.....WLKQKNKSNFVGNSTWSPY 595

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1867569 seqs, 417829326 residues

Total number of hits satisfying chosen parameters: 1867569

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications AA Main:
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2: /cgn2_6/prodata/1/pubpaa/US08_PUBCOMB.pap:*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	3231	100.0	595	US-10-659-000-4	Sequence 4, Appli
2	3231	100.0	805	US-09-978-285-2	Sequence 2, Appli
3	3231	100.0	805	US-09-999-781-2	Sequence 2, Appli
4	3231	100.0	805	US-10-158-847-142	Sequence 142, App
5	3231	100.0	805	US-10-005-956-570	Sequence 570, App
6	3231	100.0	805	US-10-005-956-843	Sequence 843, App
7	3231	100.0	805	US-10-158-825-142	Sequence 142, App
8	3231	100.0	805	US-10-158-825-142	Sequence 142, App
9	3231	100.0	805	US-10-756-149-5456	Sequence 5456, Ap
10	3231	100.0	805	US-11-059-218-2	Sequence 2, Appli
11	3231	100.0	805	US-11-059-218-106	Sequence 106, App
12	3227	99.9	805	US-10-114-893-86	Sequence 86, Appl
13	2987	92.4	681	US-09-969-384-25	Sequence 25, Appl
14	2987	92.4	681	US-10-158-847-140	Sequence 140, App
15	2987	92.4	681	US-10-158-825-140	Sequence 140, App
16	2987	92.4	681	US-10-158-825-140	Sequence 140, App
17	2987	92.4	711	US-09-969-384-13	Sequence 13, Appl
18	2987	92.4	711	US-10-158-847-138	Sequence 138, App
19	2987	92.4	711	US-10-158-825-138	Sequence 138, App
20	2987	92.4	711	US-10-158-825-138	Sequence 138, App
21	2897	89.7	555	US-10-028-072-72	Sequence 72, Appl
22	2897	89.7	555	US-10-140-808-72	Sequence 72, Appl
23	2897	89.7	555	US-10-121-049-72	Sequence 72, Appl
24	2897	89.7	555	US-10-123-904-72	Sequence 72, Appl
25	2897	89.7	555	US-10-140-470-72	Sequence 72, Appl
26	2897	89.7	555	US-10-175-746-72	Sequence 72, Appl
27	2897	89.7	555	US-10-176-918-72	Sequence 72, Appl

28	2897	89.7	555	4	US-10-176-921-72	Sequence 72, Appl
29	2897	89.7	555	4	US-10-137-865-72	Sequence 72, Appl
30	2897	89.7	555	4	US-10-140-474-72	Sequence 72, Appl
31	2897	89.7	555	4	US-10-142-431-72	Sequence 72, Appl
32	2897	89.7	555	4	US-10-143-114-72	Sequence 72, Appl
33	2897	89.7	555	4	US-10-142-419-72	Sequence 72, Appl
34	2897	89.7	555	4	US-10-123-262-72	Sequence 72, Appl
35	2897	89.7	555	4	US-10-142-423-72	Sequence 72, Appl
36	2897	89.7	555	4	US-10-121-050-72	Sequence 72, Appl
37	2897	89.7	555	4	US-10-141-755-72	Sequence 72, Appl
38	2897	89.7	555	4	US-10-143-032-72	Sequence 72, Appl
39	2897	89.7	555	4	US-10-123-108-72	Sequence 72, Appl
40	2897	89.7	555	4	US-10-123-236-72	Sequence 72, Appl
41	2897	89.7	555	4	US-10-123-261-72	Sequence 72, Appl
42	2897	89.7	555	4	US-10-140-921-72	Sequence 72, Appl
43	2897	89.7	555	4	US-10-140-928-72	Sequence 72, Appl
44	2897	89.7	555	4	US-10-121-045-72	Sequence 72, Appl
45	2897	89.7	555	4	US-10-123-292-72	Sequence 72, Appl

ALIGNMENTS

RESULT 1

US-10-659-000-4
; Sequence 4, Application US/10659000
; Publication No. US20040209344A1
; GENERAL INFORMATION:
; APPLICANT: PANTOLIANO, MICHAEL W.
; APPLICANT: RYAN, M. DOMINIC
; APPLICANT: STAKER, BART LEE
; APPLICANT: PRASAD, G. SRIDHAR
; APPLICANT: TANG, JIN
; APPLICANT: MENON, SAURABH PRABHAKAR
; APPLICANT: TOWLER, PAUL S.
; APPLICANT: WILLIAMS, DAVID H.
; APPLICANT: FISHER, MARTIN
; TITLE OF INVENTION: CRYSTAL STRUCTURE OF ANGIOTENSIN-CONVERTING ENZYME-RELATED
; FILE OF INVENTION: CARBOXYPEPTIDASE
; FILE REFERENCE: MNM/002
; CURRENT APPLICATION NUMBER: US/10/659,000
; CURRENT FILING DATE: 2003-09-09
; PRIOR APPLICATION NUMBER: 60/410,010
; PRIOR FILING DATE: 2002-09-09
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: PatentIn Ver. 3.2
; SEQ ID NO 4
; LENGTH: 595
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-659-000-4

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Best Local Similarity 100.0%; Pred. No. 1.5e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy	1	STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNTEENVQNMNAGDKWSAFLEKQST 60		
Db	1	STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNTEENVQNMNAGDKWSAFLEKQST 60		
Qy	61	LAQMPLOEIQNLTKVLQLOALQNGSSVLSEDKSKRLNTILNTMTSTYTGKVCNPDP 120		
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Qy	121	QECLLLEPGLMEIMANSLDYNERLMAWESWSEVQKLRPLYEEVVLKNEARAHYED 180		
Db	121	QECLLLEPGLMEIMANSLDYNERLMAWESWSEVQKLRPLYEEVVLKNEARAHYED 180		
Qy	181	YGDYWRGDEYNGVDGYDSRGQLIEDVEHTFEETKPLYEHLHAYVRAKLMNAPSYISP 240		
Db	181	YGDYWRGDEYNGVDGYDSRGQLIEDVEHTFEETKPLYEHLHAYVRAKLMNAPSYISP 240		
Qy	241	IGCLPAHLIGDMWGRFNTNLSLTVPFQCKPNIDVTDAWDQAWDAQRIKFAEKFPVSV 300		

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Db 241 IGCLPAHLGLDGMWGRFTWNLVSLTFPGQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSV 300
Qy 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGDRIILMCTKVWDDFLTAHHEMVG 360
Db 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGDRIILMCTKVWDDFLTAHHEMVG 360
Qy 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 420
Db 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 420
Qy 421 LKQALTIIVGTLPTFTYMLEKRWMMVFKEI PKDQWMMKWKWEMKREIIVGVVPEVPHDETYC 480
Db 421 LKQALTIIVGTLPTFTYMLEKRWMMVFKEI PKDQWMMKWKWEMKREIIVGVVPEVPHDETYC 480
Qy 481 DPASLFHVSNDYSFIRYRTLYQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 540
Db 481 DPASLFHVSNDYSFIRYRTLYQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 540
Qy 541 RLKSEPTWLALENVVGAKNMVRPLNLYFEPFLFTWLKDQKNKNSFVGWSTWSPY 595
Db 541 RLKSEPTWLALENVVGAKNMVRPLNLYFEPFLFTWLKDQKNKNSFVGWSTWSPY 595
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RESULT 2
US-09-978-385-2
; Sequence 2, Application US/09978385
; Patent No. US20020177211A1
; GENERAL INFORMATION:
; APPLICANT: Piddington, Christopher S.
; APPLICANT: Petrie, Charles
; APPLICANT: Shoemaker, Kimberly E.
; APPLICANT: Bishop, Paul D.
; TITLE OF INVENTION: ZACE2: A HUMAN METALLOENZYME
; FILE REFERENCE: 99-24C1
; CURRENT APPLICATION NUMBER: US/09/978,385
; PRIOR FILING DATE: 2001-10-16
; PRIOR APPLICATION NUMBER: 60/133,952
; PRIOR FILING DATE: 1999-05-13
; PRIOR APPLICATION NUMBER: 60/151,181
; PRIOR FILING DATE: 1999-08-27
; PRIOR APPLICATION NUMBER: 09/563,516
; PRIOR FILING DATE: 2000-05-03
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-978-385-2
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Query Match 100.0%; Score 3231; DB 3; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLKEQST 60
Db 19 STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLKEQST 78
Qy 61 LAQMPLOEQIQLTVKLOLQALQONGSSVLSBDKSKRLNTILNTMTSTIYSTGKVCNPNP 120
Db 79 LAQMPLOEQIQLTVKLOLQALQONGSSVLSBDKSKRLNTILNTMTSTIYSTGKVCNPNP 138
Qy 121 QECLLLEPGLNEIMANSLDYNERLWAWESWSEVKGQRLPLYEEYVVLKNEMARANHYED 180
Db 139 QECLLLEPGLNEIMANSLDYNERLWAWESWSEVKGQRLPLYEEYVVLKNEMARANHYED 198
Qy 181 YGDYWRGDYEVNGVDGYDSRQGLIEDVHTFEETKPLYEHLHAYVRAKLMNAYPSYISP 240
Db 199 YGDYWRGDYEVNGVDGYDSRQGLIEDVHTFEETKPLYEHLHAYVRAKLMNAYPSYISP 258
Qy 241 IGCLPAHLGLDGMWGRFTWNLVSLTFPGQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSV 300
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Db 259 IGCLPAHLGLDGMWGRFTWNLVSLTFPGQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSV 318
Qy 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGDRIILMCTKVWDDFLTAHHEMVG 360
Db 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGDRIILMCTKVWDDFLTAHHEMVG 378
Qy 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 420
Db 379 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 438
Qy 421 LKQALTIIVGTLPTFTYMLEKRWMMVFKEI PKDQWMMKWKWEMKREIIVGVVPEVPHDETYC 480
Db 439 LKQALTIIVGTLPTFTYMLEKRWMMVFKEI PKDQWMMKWKWEMKREIIVGVVPEVPHDETYC 498
Qy 481 DPASLFHVSNDYSFIRYRTLYQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 540
Db 499 DPASLFHVSNDYSFIRYRTLYQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 558
Qy 541 RLKSEPTWLALENVVGAKNMVRPLNLYFEPFLFTWLKDQKNKNSFVGWSTWSPY 595
Db 559 RLKSEPTWLALENVVGAKNMVRPLNLYFEPFLFTWLKDQKNKNSFVGWSTWSPY 613
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RESULT 3
US-09-999-781-2
; Sequence 2, Application US/09999781
; Publication No. US20040082496A1
; GENERAL INFORMATION:
; APPLICANT: ACTON, SUSAN L.
; APPLICANT: OCAIN, TIMOTHY D.
; APPLICANT: GOULD, ALEXANDRA
; APPLICANT: DALES, NATALIE A.
; APPLICANT: GUAN, BING
; APPLICANT: BROWN, JAMES A.
; APPLICANT: PATANE, MICHAEL
; APPLICANT: KADAMBI, VIVEK J.
; APPLICANT: SOLOMON, MICHAEL
; APPLICANT: STRICKER-KRONGRAD, ALAIN
; TITLE OF INVENTION: ACE-2 MODULATING COMPOUNDS AND METHODS
; TITLE OF INVENTION: OF USE THEREOF
; FILE REFERENCE: MFI-082CF4
; CURRENT APPLICATION NUMBER: US/09/999,781
; CURRENT FILING DATE: 2001-10-31
; PRIOR APPLICATION NUMBER: 09/870,382
; PRIOR FILING DATE: 2001-05-29
; PRIOR APPLICATION NUMBER: 09/704,216
; PRIOR FILING DATE: 2000-11-01
; PRIOR APPLICATION NUMBER: 60/XXX,XXX
; PRIOR FILING DATE: 2001-10-19
; NUMBER OF SEQ ID NOS: 34
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-999-781-2
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Query Match 100.0%; Score 3231; DB 3; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLKEQST 60
Db 19 STIEEQAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKWSAFLKEQST 78
Qy 61 LAQMPLOEQIQLTVKLOLQALQONGSSVLSBDKSKRLNTILNTMTSTIYSTGKVCNPNP 120
Db 79 LAQMPLOEQIQLTVKLOLQALQONGSSVLSBDKSKRLNTILNTMTSTIYSTGKVCNPNP 138
Qy 121 QECLLLEPGLNEIMANSLDYNERLWAWESWSEVKGQRLPLYEEYVVLKNEMARANHYED 180
Db 139 QECLLLEPGLNEIMANSLDYNERLWAWESWSEVKGQRLPLYEEYVVLKNEMARANHYED 198
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QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTPEETKPLYEHLHAYVRAKLMNAYPSYISLP 240
DB 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTPEETKPLYEHLHAYVRAKLMNAYPSYISLP 258
QY 241 IGCLPAHLGDMWGRFWTNLYSLTVPPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 300
DB 259 IGCLPAHLGDMWGRFWTNLYSLTVPPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMGGH 360
DB 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMGGH 378
QY 361 IOYDMAYAAQPFLLRNGANEGFHEAVGEIWSLSAATPKHLKSIIGLLSPDFQSDNTEINFP 420
DB 379 IOYDMAYAAQPFLLRNGANEGFHEAVGEIWSLSAATPKHLKSIIGLLSPDFQSDNTEINFP 438
QY 421 LKQALTIIVGTLPTFTYMLEKRWMPKGEIPKQDMKKWEMKREIVGVVBPVPHDETYC 480
DB 439 LKQALTIIVGTLPTFTYMLEKRWMPKGEIPKQDMKKWEMKREIVGVVBPVPHDETYC 498
QY 481 DPASLPHVSNDSYFIRYTRTYLQYQFOQALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 540
DB 499 DPASLPHVSNDSYFIRYTRTYLQYQFOQALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 558
QY 541 RLKSEPTWTLAENVVGAKNMVRPLNYPFPLFTWLKQDNKNSFVGWSTWSPY 595
DB 559 RLKSEPTWTLAENVVGAKNMVRPLNYPFPLFTWLKQDNKNSFVGWSTWSPY 613

RESULT 4

US-10-158-847-142
; Sequence 142, Application US/10158847
; Publication No. US20030091557A1
; GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACB-2 Activity
; FILE REFERENCE: P2557
; CURRENT APPLICATION NUMBER: US/10/158,847
; PRIOR FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/295,004
; PRIOR FILING DATE: 2001-06-04
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 142
; LENGTH: 805
; TYPE: PRT
; ORGANISM: homo sapiens
US-10-158-847-142

Query Match 100.0%; Score 3231; DB 4; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEBQAKTFLDKFNHEADLFYQSSLASWYNTNITEENVQNMNAGDKWSAFLEKEOST 60
DB 19 STIEBQAKTFLDKFNHEADLFYQSSLASWYNTNITEENVQNMNAGDKWSAFLEKEOST 78
QY 61 LAQMYPLOEIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 120
DB 79 LAQMYPLOEIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEANRANHYED 180
DB 139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEANRANHYED 198
QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTPEETKPLYEHLHAYVRAKLMNAYPSYISLP 240
DB 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTPEETKPLYEHLHAYVRAKLMNAYPSYISLP 258
QY 241 IGCLPAHLGDMWGRFWTNLYSLTVPPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 300
DB 259 IGCLPAHLGDMWGRFWTNLYSLTVPPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 318

QY 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMGGH 360
DB 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMGGH 378
QY 361 IOYDMAYAAQPFLLRNGANEGFHEAVGEIWSLSAATPKHLKSIIGLLSPDFQSDNTEINFP 420
DB 379 IOYDMAYAAQPFLLRNGANEGFHEAVGEIWSLSAATPKHLKSIIGLLSPDFQSDNTEINFP 438
QY 421 LKQALTIIVGTLPTFTYMLEKRWMPKGEIPKQDMKKWEMKREIVGVVBPVPHDETYC 480
DB 439 LKQALTIIVGTLPTFTYMLEKRWMPKGEIPKQDMKKWEMKREIVGVVBPVPHDETYC 498
QY 481 DPASLPHVSNDSYFIRYTRTYLQYQFOQALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 540
DB 499 DPASLPHVSNDSYFIRYTRTYLQYQFOQALCOAAKHEGPHLKCDISNSTEAGQKLFNNML 558
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DB 559 RLKSEPTWTLAENVVGAKNMVRPLNYPFPLFTWLKQDNKNSFVGWSTWSPY 613

RESULT 5

US-10-005-956-570
; Sequence 570, Application US/10005956
; Publication No. US20030113726A1
; GENERAL INFORMATION:
; APPLICANT: Bristol-Myers Squibb Company
; TITLE OF INVENTION: HUMAN SINGLE NUCLEOTIDE POLYMORPHISMS
; FILE REFERENCE: D0053NP
; CURRENT APPLICATION NUMBER: US/10/005,956
; CURRENT FILING DATE: 2001-12-03
; PRIOR APPLICATION NUMBER: 60/251,015
; PRIOR FILING DATE: 2000-12-04
; PRIOR APPLICATION NUMBER: 60/263,678
; PRIOR FILING DATE: 2001-01-23
; PRIOR APPLICATION NUMBER: 60/273,037
; PRIOR FILING DATE: 2001-03-02
; NUMBER OF SEQ ID NOS: 1579
; SOFTWARE: Patent in version 3.0
; SEQ ID NO 570
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-005-956-570

Query Match 100.0%; Score 3231; DB 4; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEBQAKTFLDKFNHEADLFYQSSLASWYNTNITEENVQNMNAGDKWSAFLEKEOST 60
DB 19 STIEBQAKTFLDKFNHEADLFYQSSLASWYNTNITEENVQNMNAGDKWSAFLEKEOST 78
QY 61 LAQMYPLOEIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 120
DB 79 LAQMYPLOEIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEANRANHYED 180
DB 139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEANRANHYED 198
QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTPEETKPLYEHLHAYVRAKLMNAYPSYISLP 240
DB 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTPEETKPLYEHLHAYVRAKLMNAYPSYISLP 258
QY 241 IGCLPAHLGDMWGRFWTNLYSLTVPPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 300
DB 259 IGCLPAHLGDMWGRFWTNLYSLTVPPGQKPNIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMGGH 360
DB 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRIILMCTKVTTWDDFLTAHHEMGGH 378

QY 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
DB 379 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438
QY 421 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 480
DB 439 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 498
QY 481 DPASLFHVSNDSYFIRYTRTYLQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 540
DB 499 DPASLFHVSNDSYFIRYTRTYLQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 558
QY 541 RLKSEPTWTLALENVVGAKNMVRLPLNYFEPLFTWLKDQNKNSFVGWSTWSPY 595
DB 559 RLKSEPTWTLALENVVGAKNMVRLPLNYFEPLFTWLKDQNKNSFVGWSTWSPY 613

RESULT 6

US-10-005-956-843
; Sequence 843, Application US/10005956
; Publication No. US20030113726A1
; GENERAL INFORMATION:
; APPLICANT: Bristol-Myers Squibb Company
; TITLE OF INVENTION: HUMAN SINGLE NUCLEOTIDE POLYMORPHISMS
; FILE REFERENCE: D0053NP
; CURRENT APPLICATION NUMBER: US/10/005,956
; CURRENT FILING DATE: 2001-12-03
; PRIOR APPLICATION NUMBER: 60/251,015
; PRIOR FILING DATE: 2000-12-04
; PRIOR APPLICATION NUMBER: 60/263,678
; PRIOR FILING DATE: 2001-01-23
; PRIOR APPLICATION NUMBER: 60/273,037
; PRIOR FILING DATE: 2001-03-02
; NUMBER OF SEQ ID NOS: 1579
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 843
; LENGTH: 805
; TYPE: PRT
; ORGANISM: homo sapiens
US-10-005-956-843

Query Match 100.0%; Score 3231; DB 4; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAKTFLDKFNHEAEDLFYQSSLASWYNTNITEENVQNMNAGDKWSAFLKEOST 60
DB 19 STIEEQAKTFLDKFNHEAEDLFYQSSLASWYNTNITEENVQNMNAGDKWSAFLKEOST 78
QY 61 LAQMYPLQEIQNLTVKLOLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 120
DB 79 LAQMYPLQEIQNLTVKLOLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWSEVGKQLRPLYEEYVVLKNEMARANHYED 180
DB 139 QECLLLEPGLNEIMANSLDYNERLWAWESWSEVGKQLRPLYEEYVVLKNEMARANHYED 198
QY 181 YGDYWRGDEYVNGVDGYDSRGQLTEDVEHTFEETKPLYEHLHAYVRAKLMNAYPSYISP 240
DB 199 YGDYWRGDEYVNGVDGYDSRGQLTEDVEHTFEETKPLYEHLHAYVRAKLMNAYPSYISP 258
QY 241 IGCPLPAHLGDMWGRFTWNLVSLVTPFGQKPNIDVTDAWDQAWDAQRI FKEAEKFFVSV 300
DB 259 IGCPLPAHLGDMWGRFTWNLVSLVTPFGQKPNIDVTDAWDQAWDAQRI FKEAEKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRI LMCCTKVTDWDDFLTAAHEMGH 360
DB 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRI LMCCTKVTDWDDFLTAAHEMGH 378
QY 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
DB 379 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438
QY 421 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 480
DB 439 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 498
QY 481 DPASLFHVSNDSYFIRYTRTYLQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 540
DB 499 DPASLFHVSNDSYFIRYTRTYLQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 558

QY 421 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 480
DB 439 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 498
QY 481 DPASLFHVSNDSYFIRYTRTYLQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 540
DB 499 DPASLFHVSNDSYFIRYTRTYLQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 558
QY 541 RLKSEPTWTLALENVVGAKNMVRLPLNYFEPLFTWLKDQNKNSFVGWSTWSPY 595
DB 559 RLKSEPTWTLALENVVGAKNMVRLPLNYFEPLFTWLKDQNKNSFVGWSTWSPY 613

RESULT 7

US-10-158-825-142
; Sequence 142, Application US/10158825
; Publication No. US20030138894A1
; GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACB-2 Activity
; FILE REFERENCE: PF555
; CURRENT APPLICATION NUMBER: US/10/158,825
; CURRENT FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/294,976
; PRIOR FILING DATE: 2001-06-04
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 142
; LENGTH: 805
; TYPE: PRT
; ORGANISM: homo sapiens
US-10-158-825-142

Query Match 100.0%; Score 3231; DB 4; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAKTFLDKFNHEAEDLFYQSSLASWYNTNITEENVQNMNAGDKWSAFLKEOST 60
DB 19 STIEEQAKTFLDKFNHEAEDLFYQSSLASWYNTNITEENVQNMNAGDKWSAFLKEOST 78
QY 61 LAQMYPLQEIQNLTVKLOLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 120
DB 79 LAQMYPLQEIQNLTVKLOLQALQONGSSVLSBDKSKRLNTILNTWSTIYSTGKVCNPNP 138
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWSEVGKQLRPLYEEYVVLKNEMARANHYED 180
DB 139 QECLLLEPGLNEIMANSLDYNERLWAWESWSEVGKQLRPLYEEYVVLKNEMARANHYED 198
QY 181 YGDYWRGDEYVNGVDGYDSRGQLTEDVEHTFEETKPLYEHLHAYVRAKLMNAYPSYISP 240
DB 199 YGDYWRGDEYVNGVDGYDSRGQLTEDVEHTFEETKPLYEHLHAYVRAKLMNAYPSYISP 258
QY 241 IGCPLPAHLGDMWGRFTWNLVSLVTPFGQKPNIDVTDAWDQAWDAQRI FKEAEKFFVSV 300
DB 259 IGCPLPAHLGDMWGRFTWNLVSLVTPFGQKPNIDVTDAWDQAWDAQRI FKEAEKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRI LMCCTKVTDWDDFLTAAHEMGH 360
DB 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRI LMCCTKVTDWDDFLTAAHEMGH 378
QY 361 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
DB 379 IQYDMAYAAQPFLLRNGANEGHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438
QY 421 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 480
DB 439 LLKQALTIIVGTLPTFTYMLEKRWMMVFKGEIPKQDMKKWEMKREIVGVVPEVPHDETYC 498
QY 481 DPASLFHVSNDSYFIRYTRTYLQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 540
DB 499 DPASLFHVSNDSYFIRYTRTYLQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 558

QY 541 RLKSEPTWLTALENVVGAKNMVRPLLYPEPLFTWLKDQNKNSFVGMSTWSPY 595
Db 559 RLKSEPTWLTALENVVGAKNMVRPLLYPEPLFTWLKDQNKNSFVGMSTWSPY 613

RESULT 8

US-10-158-825-142
; Sequence 142, Application US/10158825
; Publication No. US20040121429A9
; GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACE-2 Activity
; FILE REFERENCE: PF555
; CURRENT APPLICATION NUMBER: US/10/158,825
; PRIOR FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/294,976
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 142
; LENGTH: 805
; TYPE: PRT
; ORGANISM: homo sapiens
US-10-158-825-142

Query Match 100.0%; Score 3231; DB 4; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAKTFLDKFNHEADLIFYQSSLASWNTNTITEENVQNMNAGDKWSAFLKEQST 60
Db 19 STIEEQAKTFLDKFNHEADLIFYQSSLASWNTNTITEENVQNMNAGDKWSAFLKEQST 78

QY 61 LAQMYPLQBIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPDNP 120
Db 79 LAQMYPLQBIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPDNP 138

QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEMARANHYED 180
Db 139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEMARANHYED 198

QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTPEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
Db 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTPEEIKPLYEHLHAYVRAKLMNAYPSYISP 258

QY 241 IGCPLPAHLIGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 300
Db 259 IGCPLPAHLIGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 318

QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKDFRILMCTKVTDMDFLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKDFRILMCTKVTDMDFLTAHHEMGH 378

QY 361 IQYDMAYAAQPFLLRNGANEGHEAVGEMSLSAATPKHLKSIIGLLSPDPQEDNETEINF 420
Db 379 IQYDMAYAAQPFLLRNGANEGHEAVGEMSLSAATPKHLKSIIGLLSPDPQEDNETEINF 438

QY 421 LKQALTIIVGTLPFTYMLEKRWMMVFKEIIPKQDMKKWEMKREIVGVVPEVPHDETYC 480
Db 439 LKQALTIIVGTLPFTYMLEKRWMMVFKEIIPKQDMKKWEMKREIVGVVPEVPHDETYC 498

QY 481 DPASLPHVSNDSFYIRYTRTLYQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNNL 540
Db 499 DPASLPHVSNDSFYIRYTRTLYQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNNL 558

QY 541 RLKSEPTWLTALENVVGAKNMVRPLLYPEPLFTWLKDQNKNSFVGMSTWSPY 595
Db 559 RLKSEPTWLTALENVVGAKNMVRPLLYPEPLFTWLKDQNKNSFVGMSTWSPY 613

RESULT 9

US-10-756-149-5456
; Sequence 5456, Application US/10756149

; Publication No. US20050181375A1
; GENERAL INFORMATION:
; APPLICANT: Aziz, Natasha
; APPLICANT: Zlotnik, Albert
; TITLE OF INVENTION: NOVEL METHODS OF DIAGNOSIS OF METASTATIC CANCER, COMPOSITIONS AND
; TITLE OF INVENTION: METHODS OF SCREENING FOR MODULATORS OF METASTATIC CANCER
; FILE REFERENCE: file
; CURRENT APPLICATION NUMBER: US/10/756,149
; CURRENT FILING DATE: 2004-01-12
; NUMBER OF SEQ ID NOS: 5818
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 5456
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo Sapiens
US-10-756-149-5456

Query Match 100.0%; Score 3231; DB 5; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAKTFLDKFNHEADLIFYQSSLASWNTNTITEENVQNMNAGDKWSAFLKEQST 60
Db 19 STIEEQAKTFLDKFNHEADLIFYQSSLASWNTNTITEENVQNMNAGDKWSAFLKEQST 78

QY 61 LAQMYPLQBIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPDNP 120
Db 79 LAQMYPLQBIQNLTVKQLQALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPDNP 138

QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEMARANHYED 180
Db 139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVKGQLRPLYEYVVLKNEMARANHYED 198

QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTPEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
Db 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTPEEIKPLYEHLHAYVRAKLMNAYPSYISP 258

QY 241 IGCPLPAHLIGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 300
Db 259 IGCPLPAHLIGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIFKEAEKFFVSV 318

QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKDFRILMCTKVTDMDFLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKDFRILMCTKVTDMDFLTAHHEMGH 378

QY 361 IQYDMAYAAQPFLLRNGANEGHEAVGEMSLSAATPKHLKSIIGLLSPDPQEDNETEINF 420
Db 379 IQYDMAYAAQPFLLRNGANEGHEAVGEMSLSAATPKHLKSIIGLLSPDPQEDNETEINF 438

QY 421 LKQALTIIVGTLPFTYMLEKRWMMVFKEIIPKQDMKKWEMKREIVGVVPEVPHDETYC 480
Db 439 LKQALTIIVGTLPFTYMLEKRWMMVFKEIIPKQDMKKWEMKREIVGVVPEVPHDETYC 498

QY 481 DPASLPHVSNDSFYIRYTRTLYQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNNL 540
Db 499 DPASLPHVSNDSFYIRYTRTLYQFOFQALCOAAKHGEPHLKCDISNSTEAGQKLFNNL 558

QY 541 RLKSEPTWLTALENVVGAKNMVRPLLYPEPLFTWLKDQNKNSFVGMSTWSPY 595
Db 559 RLKSEPTWLTALENVVGAKNMVRPLLYPEPLFTWLKDQNKNSFVGMSTWSPY 613

RESULT 10

US-11-059-218-2
; Sequence 2, Application US/11059218
; Publication No. US20050147600A1
; GENERAL INFORMATION:
; APPLICANT: Acton, Susan L. et al.
; TITLE OF INVENTION: ANGIOTENSIN CONVERTING ENZYME HOMOLOG AND THERAPEUTIC
; TITLE OF INVENTION: DIAGNOSTIC USES THEREFOR
; FILE REFERENCE: MNI-132C93
; CURRENT APPLICATION NUMBER: US/11/059,218
; CURRENT FILING DATE: 2005-02-16


```

; PRIOR APPLICATION NUMBER: US/09/635,501
; PRIOR FILING DATE: 2000-08-09
; PRIOR APPLICATION NUMBER: 09/407,427
; PRIOR FILING DATE: 1999-09-29
; PRIOR APPLICATION NUMBER: 09/163,648
; PRIOR FILING DATE: 1998-09-30
; PRIOR APPLICATION NUMBER: 08/989,299
; PRIOR FILING DATE: 1997-12-11
; NUMBER OF SEQ ID NOS: 107
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
US-11-059-218-2

Query Match      100.0%; Score 3231; DB 6; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAKTFLDKFNHEAEDLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLEKEQST 60
Db 19 STIEEQAKTFLDKFNHEAEDLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLEKEQST 78
QY 61 LAQMYPLQEIQNLTVKLQALQALQONGSSVLSSEKSKRLNTILNTMSTIYSTGKVCNPDNP 120
Db 79 LAQMYPLQEIQNLTVKLQALQALQONGSSVLSSEKSKRLNTILNTMSTIYSTGKVCNPDNP 138
QY 121 QECLLLEPGINEIMANSLDYNERLWAWESRSEVKGQLRPLYEEYVVLKNEMARANHYED 180
Db 139 QECLLLEPGINEIMANSLDYNERLWAWESRSEVKGQLRPLYEEYVVLKNEMARANHYED 198
QY 181 YGDYWRGDIYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYIS 240
Db 199 YGDYWRGDIYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYIS 258
QY 241 IGCLPAHLGDMGWRFTWNLYSITVFPFGQKPNIDVTDAMVDQAWDAQRIFKAEAEKFFVSV 300
Db 259 IGCLPAHLGDMGWRFTWNLYSITVFPFGQKPNIDVTDAMVDQAWDAQRIFKAEAEKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAMDGLGKDFRILMCTKVWDDFLTAAHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAMDGLGKDFRILMCTKVWDDFLTAAHEMGH 378
QY 361 IOYDMAYAQQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
Db 379 IOYDMAYAQQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438
QY 421 LLKQALTIIVGTLPFTYMLEKRWMMVFKGEIPKQNMKKWEMKREIVGVVEVPVPHDETYC 480
Db 439 LLKQALTIIVGTLPFTYMLEKRWMMVFKGEIPKQNMKKWEMKREIVGVVEVPVPHDETYC 498
QY 481 DPASLPHVSNDSFYRYRTLYQFOFQALCOAAKHEGFLHKCDISNSTEAGOKLFNNML 540
Db 499 DPASLPHVSNDSFYRYRTLYQFOFQALCOAAKHEGFLHKCDISNSTEAGOKLFNNML 558
QY 541 RLKSEPTWLTALENVVGAKNMVRPLLNYFEPFLTWLKDQKNKSVFGWSTDSWSPY 595
Db 559 RLKSEPTWLTALENVVGAKNMVRPLLNYFEPFLTWLKDQKNKSVFGWSTDSWSPY 613
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RESULT 11
US-11-059-218-106
; Sequence 106, Application US/11059218
; Publication No. US20050147600A1
; GENERAL INFORMATION:
; APPLICANT: Acton, Susan L. et al.
; TITLE OF INVENTION: ANGIOGENIN CONVERTING ENZYME HOMOLOG AND THERAPEUTIC
; FILE OF INVENTION: DIAGNOSTIC USES THEREFOR
; FILE REFERENCE: MWI-132CP3
; CURRENT APPLICATION NUMBER: US/11/059,218
; CURRENT FILING DATE: 2005-02-16
; PRIOR APPLICATION NUMBER: US/09/635,501
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; PRIOR FILING DATE: 2000-08-09
; PRIOR APPLICATION NUMBER: 09/407,427
; PRIOR FILING DATE: 1999-09-29
; PRIOR APPLICATION NUMBER: 09/163,648
; PRIOR FILING DATE: 1998-09-30
; PRIOR APPLICATION NUMBER: 08/989,299
; PRIOR FILING DATE: 1997-12-11
; NUMBER OF SEQ ID NOS: 107
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 106
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
US-11-059-218-106

Query Match      100.0%; Score 3231; DB 6; Length 805;
Best Local Similarity 100.0%; Pred. No. 2.4e-273;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQAKTFLDKFNHEAEDLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLEKEQST 60
Db 19 STIEEQAKTFLDKFNHEAEDLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLEKEQST 78
QY 61 LAQMYPLQEIQNLTVKLQALQALQONGSSVLSSEKSKRLNTILNTMSTIYSTGKVCNPDNP 120
Db 79 LAQMYPLQEIQNLTVKLQALQALQONGSSVLSSEKSKRLNTILNTMSTIYSTGKVCNPDNP 138
QY 121 QECLLLEPGINEIMANSLDYNERLWAWESRSEVKGQLRPLYEEYVVLKNEMARANHYED 180
Db 139 QECLLLEPGINEIMANSLDYNERLWAWESRSEVKGQLRPLYEEYVVLKNEMARANHYED 198
QY 181 YGDYWRGDIYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYIS 240
Db 199 YGDYWRGDIYEVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYIS 258
QY 241 IGCLPAHLGDMGWRFTWNLYSITVFPFGQKPNIDVTDAMVDQAWDAQRIFKAEAEKFFVSV 300
Db 259 IGCLPAHLGDMGWRFTWNLYSITVFPFGQKPNIDVTDAMVDQAWDAQRIFKAEAEKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAMDGLGKDFRILMCTKVWDDFLTAAHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAMDGLGKDFRILMCTKVWDDFLTAAHEMGH 378
QY 361 IOYDMAYAQQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
Db 379 IOYDMAYAQQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438
QY 421 LLKQALTIIVGTLPFTYMLEKRWMMVFKGEIPKQNMKKWEMKREIVGVVEVPVPHDETYC 480
Db 439 LLKQALTIIVGTLPFTYMLEKRWMMVFKGEIPKQNMKKWEMKREIVGVVEVPVPHDETYC 498
QY 481 DPASLPHVSNDSFYRYRTLYQFOFQALCOAAKHEGFLHKCDISNSTEAGOKLFNNML 540
Db 499 DPASLPHVSNDSFYRYRTLYQFOFQALCOAAKHEGFLHKCDISNSTEAGOKLFNNML 558
QY 541 RLKSEPTWLTALENVVGAKNMVRPLLNYFEPFLTWLKDQKNKSVFGWSTDSWSPY 595
Db 559 RLKSEPTWLTALENVVGAKNMVRPLLNYFEPFLTWLKDQKNKSVFGWSTDSWSPY 613
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RESULT 12
US-10-114-893-86
; Sequence 86, Application US/10114893
; Publication No. US20020193567A1
; GENERAL INFORMATION:
; APPLICANT: Jacobs, Kenneth
; APPLICANT: McCoy, John M.
; APPLICANT: LaVallie, Edward R.
; APPLICANT: Collins-Racie, Lisa A.
; APPLICANT: Evans, Cheryl
; APPLICANT: Merberg, David
; APPLICANT: Treacy, Maurice
; APPLICANT: Bowman, Michael R.
```

APPLICANT: Spaulding, Vikki
APPLICANT: Carlin-Duckett, McKeough
APPLICANT: Kelleher, Kerry S.
APPLICANT: Genetics Institute, Inc.
TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES ENCODING THEM
FILE REFERENCE: GI 6000-10A
CURRENT APPLICATION NUMBER: US/10/114,893
CURRENT FILING DATE: 2002-04-02
EARLIER APPLICATION NUMBER: 09/413,232
EARLIER FILING DATE: 1999-10-06
NUMBER OF SEQ ID NOS: 321
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 86
LENGTH: 805
TYPE: PRT
ORGANISM: Homo sapiens
US-10-114-893-86

Query Match 99.9%; Score 3227; DB 4; Length 805;
Best Local Similarity 99.8%; Pred. No. 5.4e-273;
Matches 594; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEAKTFLDKFHEADLDFQSSLASWNTNTNTEENVQNMNAGDKNSAFLEKQST 60
DB |||||
QY 19 STIEEAKTFLDKFHEADLDFQSSLASWNTNTNTEENVQNMNAGDKNSAFLEKQST 78
DB |||||
QY 61 LAQMYPLOBIQNLTVKQLQALQOQSSVLSBDKSKRLNTILNTWSTYTGKVCNPNP 120
DB |||||
QY 79 LAQMYPLOBIQNLTVKQLQALQOQSSVLSBDKSKRLNTILNTWSTYTGKVCNPNP 138
DB |||||
QY 121 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEYVVLKNEARANYED 180
DB |||||
QY 139 QECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYEYVVLKNEARANYED 198
DB |||||
QY 181 YGDYWRGDEYVNGVDYDSRGQLIEDVHTFEEIKPLYEHLHAYVRAKLMNAYPSYIS 240
DB |||||
QY 199 YGDYWRGDEYVNGVDYDSRGQLIEDVHTFEEIKPLYEHLHAYVRAKLMNAYPSYIS 258
DB |||||
QY 241 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIKFAEAKFFVS 300
DB |||||
QY 259 IGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDQAWDAQRIKFAEAKFFVS 318
DB |||||
QY 301 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRLMCTKVTWDDFLTAHHEMGH 360
DB |||||
QY 319 GLPNMTQGFWNSMLTDPGNQKAVCHPTAWDLGKGFRLMCTKVTWDDFLTAHHEMGH 378
DB |||||
QY 361 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 420
DB |||||
QY 379 IQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNTEINF 438
DB |||||
QY 421 LLKQALTIIVGTLPTFTYMLEKRWMPFKGEIPKQDMKKWEMKREIVGVVFPVPHDETYC 480
DB |||||
QY 439 LLKQALTIIVGTLPTFTYMLEKRWMPFKGEIPKQDMKKWEMKREIVGVVFPVPHDETYC 498
DB |||||
QY 481 DPASLPHVNSDYSFIRYTRTLTYQFQFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 540
DB |||||
QY 499 DPASLPHVNSDYSFIRYTRTLTYQFQFQALCOAAKHGEPHLKCDISNSTEAGQKLFNML 558
DB |||||
QY 541 RLKSEFPWTALLENVVGAKNMVRPLLNYFPELFTWLKDONKNSFVGWSTDWSPY 595
DB |||||
QY 559 RLKSEFPWTALLENVVGAKNMVRPLLNYFPELFTWLKDONKNSFVGWSTDWSPY 613
DB |||||

RESULT 13
US-09-969-384-25
Sequence 25, Application US/09969384
Publication No. US20020192749A1
GENERAL INFORMATION:
APPLICANT: Moore, et al.
TITLE OF INVENTION: Human Gene Polynucleotides, Polypeptides, and Antibodies
FILE REFERENCE: PT055P1
CURRENT APPLICATION NUMBER: US/09/969,384
CURRENT FILING DATE: 2001-10-03

PRIOR APPLICATION NUMBER: PCT/US01/10542
PRIOR FILING DATE: 2001-04-02
PRIOR APPLICATION NUMBER: 60/236,384
PRIOR FILING DATE: 2000-09-29
PRIOR APPLICATION NUMBER: 60/194,118
PRIOR FILING DATE: 2000-04-03
NUMBER OF SEQ ID NOS: 27
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 25
LENGTH: 681
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: SITE
LOCATION: (219)
OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
NAME/KEY: SITE
LOCATION: (240)
OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
NAME/KEY: SITE
LOCATION: (499)
OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
US-09-969-384-25

Query Match 92.4%; Score 2987; DB 3; Length 681;
Best Local Similarity 99.3%; Pred. No. 4.3e-252;
Matches 548; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 44 MNAGDKNSAFLEKQSTLAQMYPLOBIQNLTVKQLQALQOQSSVLSBDKSKRLNTILN 103
DB |||||
QY 1 MNAGDKNSAFLEKQSTLAQMYPLOBIQNLTVKQLQALQOQSSVLSBDKSKRLNTILN 60
DB |||||
QY 104 TWSTYTGKVCNPNPQECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYE 163
DB |||||
QY 61 TWSTYTGKVCNPNPQECLLLEPGLNEIMANSLDYNERLWAWESWRSEVGKQLRPLYE 120
DB |||||
QY 164 EYVVLKNEARANYEDYGDYWRGDEYVNGVDYDSRGQLIEDVHTFEEIKPLYEHLH 223
DB |||||
QY 121 EYVVLKNEARANYEDYGDYWRGDEYVNGVDYDSRGQLIEDVHTFEEIKPLYEHLH 180
DB |||||
QY 224 AYVRAKLMNAYPSYISPIGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDOA 283
DB |||||
QY 181 AYVRAKLMNAYPSYISPIGCLPAHLGDMWGRFWTNLYSLTVPFGQKPNIDVTDAMVDOX 240
DB |||||
QY 284 WDAQRIKFAEAKFFVSUGLNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMCM 343
DB |||||
QY 241 WDAQRIKFAEAKFFVSUGLNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMCM 300
DB |||||
QY 344 TKVTWDDFLTAHHEMGHITQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSI 403
DB |||||
QY 301 TKVTWDDFLTAHHEMGHITQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSI 360
DB |||||
QY 404 GLLSPDFQEDNTEINFLLKQALTIIVGTLPTFTYMLEKRWMPFKGEIPKQDMKKWEMK 463
DB |||||
QY 361 GLLSPDFQEDNTEINFLLKQALTIIVGTLPTFTYMLEKRWMPFKGEIPKQDMKKWEMK 420
DB |||||
QY 464 REIVGVVFPVPHDETYCDPASILFHVNSDYSFIRYTRTLTYQFQFQALCOAAKHGEPHLK 523
DB |||||
QY 421 REIVGVVFPVPHDETYCDPASILFHVNSDYSFIRYTRTLTYQFQFQALCOAAKHGEPHLK 480
DB |||||
QY 524 CDISNSTEAGQKLFNMLRLKSEFPWTALLENVVGAKNMVRPLLNYFPELFTWLKDONKN 583
DB |||||
QY 481 CDISNSTEAGQKLFNMLRXGKSEFPWTALLENVVGAKNMVRPLLNYFPELFTWLKDONKN 540
DB |||||
QY 584 SFVGWSTDWSPY 595
DB |||||
QY 541 SFVGWSTDWSPY 552
DB |||||

RESULT 14
US-10-158-847-140
Sequence 140, Application US/10158847
Publication No. US20030091557A1

GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACE-2 Activity
; FILE REFERENCE: PF557
; CURRENT APPLICATION NUMBER: US/10/158,847
; PRIOR FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/295,004
; PRIOR FILING DATE: 2001-06-04
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 140
; LENGTH: 681
; TYPE: PRT
; ORGANISM: homo sapiens
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (219)..(219)
; OTHER INFORMATION: Xaa equals any amino acid
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (240)..(240)
; OTHER INFORMATION: Xaa equals any amino acid
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (499)..(499)
; OTHER INFORMATION: Xaa equals any amino acid
US-10-158-847-140

Query Match 92.4%; Score 2987; DB 4; Length 681;
Best Local Similarity 99.3%; Pred. No. 4.3e-252;
Matches 548; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY	44	MNAGDKSAFLKEQSTLAQMYPLOEIQNLTVKQLQALQOQSSVLSSEKSKRLNTILN	103
DB	1	MNAGDKSAFLKEQSTLAQMYPLOEIQNLTVKQLQALQOQSSVLSSEKSKRLNTILN	60
QY	104	TMSTIYSTGKVCNPNPQECLLLEPGLNEIMANSLDYNERLWAWESRSEVGKQLRPLYE	163
DB	61	TMSTIYSTGKVCNPNPQECLLLEPGLNEIMANSLDYNERLWAWESRSEVGKQLRPLYE	120
QY	164	EYVLKNEMARANHYEDYDGYWRGDEYVNGVDYDYSRGQLIEDVEHTFEETKPLYEHLH	223
DB	121	EYVLKNEMARANHYEDYDGYWRGDEYVNGVDYDYSRGQLIEDVEHTFEETKPLYEHLH	180
QY	224	AYVRKLMNAYPSYISPIGCLPAHLGDMWGRFWTNLYSLTVPFQCKENIDVTDAMVDQA	283
DB	181	AYVRKLMNAYPSYISPIGCLPAHLGDMWGRFWTNLYSLTVPFQCKENIDVTDAMVDQA	240
QY	284	WDAQRIFKEAEKFFVSVGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMC	343
DB	241	WDAQRIFKEAEKFFVSVGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMC	300
QY	344	TKVTWDDFLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSI	403
DB	301	TKVTWDDFLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSI	360
QY	404	GLLSPDFQEDNETEINFLKQALTIIVGTLPTTYMLEKRWMMVFKEIPKQWMMKWMEMK	463
DB	361	GLLSPDFQEDNETEINFLKQALTIIVGTLPTTYMLEKRWMMVFKEIPKQWMMKWMEMK	420
QY	464	REIVGVVEVPVPHDETYCDPASLPHVSNDSYFIRYTRTLYQFQFQALCOAAKHGEGPLHK	523
DB	421	REIVGVVEVPVPHDETYCDPASLPHVSNDSYFIRYTRTLYQFQFQALCOAAKHGEGPLHK	480
QY	524	CDISNSTEAGOKLFNNLRKGKSEPTLALENVVGAQNNVRPLLNYFEPFLTWLKDQKNK	583
DB	481	CDISNSTEAGOKLFNNLRKGKSEPTLALENVVGAQNNVRPLLNYFEPFLTWLKDQKNK	540
QY	584	SPVGMSTWDSPY 595	
DB	541	SPVGMSTWDSPY 552	

RESULT 15
US-10-158-825-140
; Sequence 140, Application US/10158825
; Publication No. US20030138894A1
; GENERAL INFORMATION:
; APPLICANT: Tom Parry et al.
; TITLE OF INVENTION: Method and Compositions for Modulating ACE-2 Activity
; FILE REFERENCE: PF555
; CURRENT APPLICATION NUMBER: US/10/158,825
; CURRENT FILING DATE: 2002-06-03
; PRIOR APPLICATION NUMBER: 60/294,976
; PRIOR FILING DATE: 2001-06-04
; NUMBER OF SEQ ID NOS: 158
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 140
; LENGTH: 681
; TYPE: PRT
; ORGANISM: homo sapiens
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (219)..(219)
; OTHER INFORMATION: Xaa equals any amino acid
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (240)..(240)
; OTHER INFORMATION: Xaa equals any amino acid
; FEATURE:
; NAME/KEY: MISC FEATURE
; LOCATION: (499)..(499)
; OTHER INFORMATION: Xaa equals any amino acid
US-10-158-825-140

Query Match 92.4%; Score 2987; DB 4; Length 681;
Best Local Similarity 99.3%; Pred. No. 4.3e-252;
Matches 548; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY	44	MNAGDKSAFLKEQSTLAQMYPLOEIQNLTVKQLQALQOQSSVLSSEKSKRLNTILN	103
DB	1	MNAGDKSAFLKEQSTLAQMYPLOEIQNLTVKQLQALQOQSSVLSSEKSKRLNTILN	60
QY	104	TMSTIYSTGKVCNPNPQECLLLEPGLNEIMANSLDYNERLWAWESRSEVGKQLRPLYE	163
DB	61	TMSTIYSTGKVCNPNPQECLLLEPGLNEIMANSLDYNERLWAWESRSEVGKQLRPLYE	120
QY	164	EYVLKNEMARANHYEDYDGYWRGDEYVNGVDYDYSRGQLIEDVEHTFEETKPLYEHLH	223
DB	121	EYVLKNEMARANHYEDYDGYWRGDEYVNGVDYDYSRGQLIEDVEHTFEETKPLYEHLH	180
QY	224	AYVRKLMNAYPSYISPIGCLPAHLGDMWGRFWTNLYSLTVPFQCKENIDVTDAMVDQA	283
DB	181	AYVRKLMNAYPSYISPIGCLPAHLGDMWGRFWTNLYSLTVPFQCKENIDVTDAMVDQA	240
QY	284	WDAQRIFKEAEKFFVSVGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMC	343
DB	241	WDAQRIFKEAEKFFVSVGLPNNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMC	300
QY	344	TKVTWDDFLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSI	403
DB	301	TKVTWDDFLTAHHEMGHIQYDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSI	360
QY	404	GLLSPDFQEDNETEINFLKQALTIIVGTLPTTYMLEKRWMMVFKEIPKQWMMKWMEMK	463
DB	361	GLLSPDFQEDNETEINFLKQALTIIVGTLPTTYMLEKRWMMVFKEIPKQWMMKWMEMK	420
QY	464	REIVGVVEVPVPHDETYCDPASLPHVSNDSYFIRYTRTLYQFQFQALCOAAKHGEGPLHK	523
DB	421	REIVGVVEVPVPHDETYCDPASLPHVSNDSYFIRYTRTLYQFQFQALCOAAKHGEGPLHK	480
QY	524	CDISNSTEAGOKLFNNLRKGKSEPTLALENVVGAQNNVRPLLNYFEPFLTWLKDQKNK	583
DB	481	CDISNSTEAGOKLFNNLRKGKSEPTLALENVVGAQNNVRPLLNYFEPFLTWLKDQKNK	540
QY	584	SPVGMSTWDSPY 595	

Db 541 SFVGNSTWSPY 552
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Search completed: March 28, 2006, 11:42:32
Job time : 97.1865 secs

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OM protein - protein search, using sw model

Run on: March 28, 2006, 11:38:04 ; Search time 13.1176 Seconds
(without alignments)
1337.835 Million cell updates/sec

Title: US-10-659-000-4
Perfect score: 3231
Sequence: 1 STIEEQAKTFLDKFHAEAD.....WLKQNKNSFVGNSTDNPSY 595

Scoring table: BLOSUM62

Searched: 174695 seqs, 29494374 residues

Total number of hits satisfying chosen parameters: 174695

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

- Database : Published Applications AA New.*
- 1: /SIDSS/ptodata/1/pubppa/US08_NEW_PUB.pep.*
 - 2: /SIDSS/ptodata/1/pubppa/US06_NEW_PUB.pep.*
 - 3: /SIDSS/ptodata/1/pubppa/US07_NEW_PUB.pep.*
 - 4: /SIDSS/ptodata/1/pubppa/PCT_NEW_PUB.pep.*
 - 5: /SIDSS/ptodata/1/pubppa/US05_NEW_PUB.pep.*
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 - 7: /SIDSS/ptodata/1/pubppa/US11_NEW_PUB.pep.*
 - 8: /SIDSS/ptodata/1/pubppa/US60_NEW_PUB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	3231	100.0	805	US-10-518-599-2	Sequence 2, Appli
2	3231	100.0	805	US-10-957-880-1	Sequence 1, Appli
3	3218	99.6	702	US-10-957-880-2	Sequence 2, Appli
4	2897	89.7	555	US-10-131-826A-72	Sequence 72, Appl
5	2897	89.7	555	US-10-973-115B-72	Sequence 72, Appl
6	2757	85.3	805	US-10-518-599-24	Sequence 24, Appli
7	2755	85.3	805	US-10-518-599-4	Sequence 4, Appli
8	1335	41.3	732	US-10-518-599-23	Sequence 23, Appl
9	1335	41.3	732	US-10-995-561-1020	Sequence 1020, Ap
10	1335	41.3	1160	US-10-995-561-1019	Sequence 1019, Ap
11	1335	41.3	1302	US-10-995-561-1024	Sequence 1024, Ap
12	1335	41.3	1306	US-10-995-561-1027	Sequence 1027, Ap
13	1334	41.3	732	US-10-518-599-22	Sequence 22, Appli
14	1269	39.3	616	US-10-995-561-1018	Sequence 1018, Ap
15	1269	39.3	616	US-10-995-561-1022	Sequence 1022, Ap
16	1229	38.0	638	US-10-995-561-1025	Sequence 1025, Ap
17	1224	37.9	560	US-10-995-561-1026	Sequence 1026, Ap
18	1125	34.8	626	US-10-533-811-43	Sequence 43, Appli
19	909.5	28.1	424	US-10-995-561-1017	Sequence 1017, Ap
20	433.5	13.4	254	US-10-995-561-1021	Sequence 1021, Ap
21	266.5	8.2	209	US-10-995-561-1023	Sequence 1023, Ap
22	122	3.8	3488	US-11-087-099-9005	Sequence 9005, Ap
23	114	3.5	1493	US-10-330-773-502	Sequence 502, Appl
24	108.5	3.4	3487	US-10-330-773-502	Sequence 502, Appl
25	108	3.3	3487	US-11-087-099-9068	Sequence 9068, Ap

ALIGNMENTS

RESULT 1

US-10-518-599-2
; Sequence 2, Application US/10518599
; Publication No. US20050251873A1
; GENERAL INFORMATION:
; APPLICANT: PENNINGER, JOSEPH M.
; APPLICANT: CRACKOWER, MICHAEL A.
; TITLE OF INVENTION: ACE2 ACTIVATION FOR TREATMENT OF HEART, LUNG AND
; TITLE OF INVENTION: KIDNEY DISEASE AND HYPERTENSION
; FILE REFERENCE: SOAN:064US
; CURRENT APPLICATION NUMBER: US/10/518,599
; CURRENT FILING DATE: 2004-12-17
; PRIOR APPLICATION NUMBER: PCT/CA03/000882
; PRIOR FILING DATE: 2003-06-19
; PRIOR APPLICATION NUMBER: US 60/389,709
; PRIOR FILING DATE: 2002-06-19
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-518-599-2

Query Match	100.0%;	Score 3231;	DB 6;	Length 805;
Best Local Similarity	100.0%;	Pred. No. 1.3e-252;		
Matches 595;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
Qy	1	STIEEQAKTFLDKFNHEAEDLFYQSSLSASWNYNTNITEENVQNNNAGDKWSAFLEKQST	60	
Db	19	STIEEQAKTFLDKFNHEAEDLFYQSSLSASWNYNTNITEENVQNNNAGDKWSAFLEKQST	78	
Qy	61	LAQMPYPLQEIQTIVKLQALQONGSSVLSEDKSKRLNTILNTMTSTIYSGKVCNPNP	120	
Db	79	LAQMPYPLQEIQTIVKLQALQONGSSVLSEDKSKRLNTILNTMTSTIYSGKVCNPNP	138	
Qy	121	QECLLEPGLNETMANSLDYNERLWAWESRSEVKGQLRPLYEEYVVLKNEMARANHYYED	180	
Db	139	QECLLEPGLNETMANSLDYNERLWAWESRSEVKGQLRPLYEEYVVLKNEMARANHYYED	198	
Qy	181	YGDYWRGDFYNGVDGYDSRGQLIEDVEHTFBEIKPLYEHLHAYVRAKLNAYPSYISP	240	
Db	199	YGDYWRGDFYNGVDGYDSRGQLIEDVEHTFBEIKPLYEHLHAYVRAKLNAYPSYISP	258	
Qy	241	IGCLPAHLGLDMWGRFWTNLSYLTVPFGQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSV	300	
Db	259	IGCLPAHLGLDMWGRFWTNLSYLTVPFGQKPNIDVTDAMVDQAWDAQRIKFAEKFFVSV	318	

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QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVTMDDFLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTANDLGKGFRLMCTKVTMDDFLTAHHEMGH 378
QY 361 IQYDMAYAAQPFLLRNGANGEGHEAVGEIMSLSAATPKHLKSGILGLSPDFQSDNETEINF 420
Db 379 IQYDMAYAAQPFLLRNGANGEGHEAVGEIMSLSAATPKHLKSGILGLSPDFQSDNETEINF 438
QY 421 LKQALTIIVGTLPTFTYMLEKRWMMVKGEIPKQOMKKWEMKREIVGVVBPVPHDETYC 480
Db 439 LKQALTIIVGTLPTFTYMLEKRWMMVKGEIPKQOMKKWEMKREIVGVVBPVPHDETYC 498
QY 481 DPASLFHVSNDSYFIRYTRTYQFQEQALCOAAKHGEPHLHKCDISNSTEAGQKLFNNML 540
Db 499 DPASLFHVSNDSYFIRYTRTYQFQEQALCOAAKHGEPHLHKCDISNSTEAGQKLFNNML 558
QY 541 RLKGSPEWTLALENVVGAKNMNVRPLLNYFEPLFTWLKDQNKNSFVGWSTWDSPY 595
Db 559 RLKGSPEWTLALENVVGAKNMNVRPLLNYFEPLFTWLKDQNKNSFVGWSTWDSPY 613

RESULT 2
US-10-957-880-1
; Sequence 1, Application US/10957880
; Publication No. US20050282154A1
; GENERAL INFORMATION:
; APPLICANT: Brigham & Women's Hospital, Inc.
; APPLICANT: Farzan, Michael R
; APPLICANT: Li, Wenhui
; APPLICANT: Moore, Michael J
; TITLE OF INVENTION: Angiotensin-Converting Enzyme-2 as a Receptor for the SARS Corona
; FILE REFERENCE: 7570/80644
; CURRENT APPLICATION NUMBER: US/10/957,880
; CURRENT FILING DATE: 2004-10-05
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-957-880-1
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Query Match 100.0%; Score 3231; DB 6; Length 805;
Best Local Similarity 100.0%; Pred. No. 1.3e-252;
Matches 595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEQAKTFLDKFNHEADLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLKEQST 60
Db 19 STIEQAKTFLDKFNHEADLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLKEQST 78
QY 61 LAQMYPLOEQIONLTVKLQLOALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 120
Db 79 LAQMYPLOEQIONLTVKLQLOALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNP 138
QY 121 QECCLLEPGLNEIMANSLDYNERLWAWESWRSSEVGKQLRPLYEEYVVLKKNEMARANHVED 180
Db 139 QECCLLEPGLNEIMANSLDYNERLWAWESWRSSEVGKQLRPLYEEYVVLKKNEMARANHVED 198
QY 181 YGDYWRGDEYVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 240
Db 199 YGDYWRGDEYVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISP 258
QY 241 IGCLPAHLGDMWGRFTNLYSLTVPFGQKPNIDVTDAWVQAWDAQRIKFAEAKFFVSV 300
Db 259 IGCLPAHLGDMWGRFTNLYSLTVPFGQKPNIDVTDAWVQAWDAQRIKFAEAKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVTMDDFLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTANDLGKGFRLMCTKVTMDDFLTAHHEMGH 378
QY 361 IQYDMAYAAQPFLLRNGANGEGHEAVGEIMSLSAATPKHLKSGILGLSPDFQSDNETEINF 420
Db 379 IQYDMAYAAQPFLLRNGANGEGHEAVGEIMSLSAATPKHLKSGILGLSPDFQSDNETEINF 438
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QY 421 LKQALTIIVGTLPTFTYMLEKRWMMVKGEIPKQOMKKWEMKREIVGVVBPVPHDETYC 480
Db 439 LKQALTIIVGTLPTFTYMLEKRWMMVKGEIPKQOMKKWEMKREIVGVVBPVPHDETYC 498
QY 481 DPASLFHVSNDSYFIRYTRTYQFQEQALCOAAKHGEPHLHKCDISNSTEAGQKLFNNML 540
Db 499 DPASLFHVSNDSYFIRYTRTYQFQEQALCOAAKHGEPHLHKCDISNSTEAGQKLFNNML 558
QY 541 RLKGSPEWTLALENVVGAKNMNVRPLLNYFEPLFTWLKDQNKNSFVGWSTWDSPY 595
Db 559 RLKGSPEWTLALENVVGAKNMNVRPLLNYFEPLFTWLKDQNKNSFVGWSTWDSPY 613

RESULT 3
US-10-957-880-2
; Sequence 2, Application US/10957880
; Publication No. US20050282154A1
; GENERAL INFORMATION:
; APPLICANT: Brigham & Women's Hospital, Inc.
; APPLICANT: Farzan, Michael R
; APPLICANT: Li, Wenhui
; APPLICANT: Moore, Michael J
; TITLE OF INVENTION: Angiotensin-Converting Enzyme-2 as a Receptor for the SARS Corona
; FILE REFERENCE: 7570/80644
; CURRENT APPLICATION NUMBER: US/10/957,880
; CURRENT FILING DATE: 2004-10-05
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 702
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-957-880-2
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Query Match 99.6%; Score 3218; DB 6; Length 702;
Best Local Similarity 100.0%; Pred. No. 1.2e-251;
Matches 592; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 BEQAKTFLDKFNHEADLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLKEQSTLAQ 63
Db 1 BEQAKTFLDKFNHEADLFYQSSLASWNTNTITEENVQNMNAGDKWSAFLKEQSTLAQ 60
QY 64 MYPLQEQIONLTVKLQLOALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNPQBC 123
Db 61 MYPLQEQIONLTVKLQLOALQONGSSVLSSEKSKRLNTILNTMTSTIYSTGKVCNPNPQBC 120
QY 124 LLLLEPGLNEIMANSLDYNERLWAWESWRSSEVGKQLRPLYEEYVVLKKNEMARANHVEDYGD 183
Db 121 LLLLEPGLNEIMANSLDYNERLWAWESWRSSEVGKQLRPLYEEYVVLKKNEMARANHVEDYGD 180
QY 184 YWRGDEYVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISPIC 243
Db 181 YWRGDEYVNGVDGYDSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLMNAYPSYISPIC 240
QY 244 LPAHLGDMWGRFTNLYSLTVPFGQKPNIDVTDAWVQAWDAQRIKFAEAKFFVSVGLP 303
Db 241 LPAHLGDMWGRFTNLYSLTVPFGQKPNIDVTDAWVQAWDAQRIKFAEAKFFVSVGLP 300
QY 304 NMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVTMDDFLTAHHEMGHIOY 363
Db 301 NMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRLMCTKVTMDDFLTAHHEMGHIOY 360
QY 364 DMAYAAQPFLLRNGANGEGHEAVGEIMSLSAATPKHLKSGILGLSPDFQSDNETEINFLLK 423
Db 361 DMAYAAQPFLLRNGANGEGHEAVGEIMSLSAATPKHLKSGILGLSPDFQSDNETEINFLLK 420
QY 424 QALTIIVGTLPTFTYMLEKRWMMVKGEIPKQOMKKWEMKREIVGVVBPVPHDETYCDBA 483
Db 421 QALTIIVGTLPTFTYMLEKRWMMVKGEIPKQOMKKWEMKREIVGVVBPVPHDETYCDBA 480
QY 484 SLFHVSNDSYFIRYTRTYQFQEQALCOAAKHGEPHLHKCDISNSTEAGQKLFNNMLRIG 543
Db 481 SLFHVSNDSYFIRYTRTYQFQEQALCOAAKHGEPHLHKCDISNSTEAGQKLFNNMLRIG 540
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Db 481 SLFHVSNDYSPRYRTLYQFOFQALCOAAGHEGLHKCDISNSTEAGQKLFNMLRLG 540
QY 544 KSEPTWLALENVVGAKNNVRPLLNYPEPLFTWLKDQNNKNSFGWSTWSPY 595
Db 541 KSEPTWLALENVVGAKNNVRPLLNYFEPLFTWLKDQNNKNSFGWSTWSPY 592

RESULT 4

US-10-131-826A-72
; Sequence 72, Application US/10131826A
; Publication No. US20050245730A1
; GENERAL INFORMATION:
; APPLICANT: Baker, Kevin P.
; APPLICANT: Beresini, Maureen
; APPLICANT: Deforge, Laura
; APPLICANT: Desnoyers, Luc
; APPLICANT: Filvaroff, Ellen
; APPLICANT: Gao, Wei-Qiang
; APPLICANT: Gerritsen, Mary E.
; APPLICANT: Goddard, Audrey
; APPLICANT: Godowski, Paul
; APPLICANT: Gurney, Austin L.
; APPLICANT: Sherwood, Steven
; APPLICANT: Smith, Victoria
; APPLICANT: Stewart, Timothy A.
; APPLICANT: Tumas, Daniel
; APPLICANT: Watanabe, Colin K
; APPLICANT: Wood, William
; APPLICANT: Zhang, Zemin

; TITLE OF INVENTION: SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC
; FILE REFERENCE: P330R1C128

; CURRENT APPLICATION NUMBER: US/10/131,826A
; CURRENT FILING DATE: 2002-04-24
; PRIOR APPLICATION NUMBER: 60/049911
; PRIOR FILING DATE: 1997-06-18
; PRIOR APPLICATION NUMBER: 60/056974
; PRIOR FILING DATE: 1997-08-26
; PRIOR APPLICATION NUMBER: 60/059113
; PRIOR FILING DATE: 1997-09-17
; PRIOR APPLICATION NUMBER: 60/059115
; PRIOR FILING DATE: 1997-09-17
; PRIOR APPLICATION NUMBER: 60/059117
; PRIOR FILING DATE: 1997-09-17
; PRIOR APPLICATION NUMBER: 60/059122
; PRIOR FILING DATE: 1997-09-17
; PRIOR APPLICATION NUMBER: 60/059184
; PRIOR FILING DATE: 1997-09-17
; PRIOR APPLICATION NUMBER: 60/059263
; PRIOR FILING DATE: 1997-09-18
; PRIOR APPLICATION NUMBER: 60/059352
; PRIOR FILING DATE: 1997-09-19
; PRIOR APPLICATION NUMBER: 60/059588
; PRIOR FILING DATE: 1997-09-19
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 550
; SEQ ID NO 72
; LENGTH: 555
; TYPE: PRT
; ORGANISM: Homo Sapien

US-10-131-826A-72
Query Match 89.7%; Score 2897; DB 6; Length 555;
Best Local Similarity 99.8%; Pred. No. 6.7e-226;
Matches 535; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 STIEEQATFLDKFNHEADLFYQSSLASWNTNTIENVQNMNAGDKSAFLKEQST 60
Db 19 STIEEQATFLDKFNHEADLFYQSSLASWNTNTIENVQNMNAGDKSAFLKEQST 78
QY 61 LAQMYPLOEQIONLTVKQLQALQOQSSVLSEDKSKRLNTILNTWSTIYTGKVCNPDNP 120
Db 79 LAQMYPLOEQIONLTVKQLQALQOQSSVLSEDKSKRLNTILNTWSTIYTGKVCNPDNP 138

RESULT 5

US-10-973-115B-72

; Sequence 72, Application US/10973115B
; Publication No. US20060040351A1
; GENERAL INFORMATION:
; APPLICANT: Baker, Kevin P.
; APPLICANT: Beresini, Maureen
; APPLICANT: Deforge, Laura
; APPLICANT: Desnoyers, Luc
; APPLICANT: Filvaroff, Ellen
; APPLICANT: Gao, Wei-Qiang
; APPLICANT: Gerritsen, Mary E.
; APPLICANT: Goddard, Audrey
; APPLICANT: Godowski, Paul
; APPLICANT: Gurney, Austin L.
; APPLICANT: Sherwood, Steven
; APPLICANT: Smith, Victoria
; APPLICANT: Stewart, Timothy A.
; APPLICANT: Tumas, Daniel
; APPLICANT: Watanabe, Colin K.
; APPLICANT: Wood, William I.
; APPLICANT: Zhang, Zemin
; TITLE OF INVENTION: SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC ACIDS ENCODING
; FILE REFERENCE: 39870-3330R1C300C1
; CURRENT APPLICATION NUMBER: US/10/973,115B
; CURRENT FILING DATE: 2004-10-22
; PRIOR APPLICATION NUMBER: US 10/145,747
; PRIOR FILING DATE: 2002-05-14
; PRIOR APPLICATION NUMBER: US 10/028,072
; PRIOR FILING DATE: 2001-12-19
; PRIOR APPLICATION NUMBER: PCT/US00/32678
; PRIOR FILING DATE: 2000-12-01
; PRIOR APPLICATION NUMBER: US 09/581,742
; PRIOR FILING DATE: 2000-06-16
; PRIOR APPLICATION NUMBER: PCT/US00/05746
; PRIOR FILING DATE: 2000-03-02
; PRIOR APPLICATION NUMBER: US 60/135,736
; PRIOR FILING DATE: 1999-05-25
; PRIOR APPLICATION NUMBER: US 60/123,090
; PRIOR FILING DATE: 1999-03-05
; NUMBER OF SEQ ID NOS: 550
; SEQ ID NO 72

QY 121 QECLLLEPLCNEIMANSIDYNERLWAMESWSEVKGQKRLPLYEEYVVLKNEKMARANHYED 180
Db 139 QECLLLEPLCNEIMANSIDYNERLWAMESWSEVKGQKRLPLYEEYVVLKNEKMARANHYED 198
QY 181 YGDYWRGDIYVNGVDYDYSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLNAYPSYISP 240
Db 199 YGDYWRGDIYVNGVDYDYSRGQLIEDVEHTFEEIKPLYEHLHAYVRAKLNAYPSYISP 258
QY 241 IGCLPAHLGLDMGRFWTNLYSLTVPFQCKENIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 300
Db 259 IGCLPAHLGLDMGRFWTNLYSLTVPFQCKENIDVTDAMVDQAWDAQRIKFAEAEKFFVSV 318
QY 301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTADNLGKDFRIILMCTKVTDWDDPLTAHHEMGH 360
Db 319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTADNLGKDFRIILMCTKVTDWDDPLTAHHEMGH 378
QY 361 IQDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
Db 379 IQDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438
QY 421 LLKQALTTVGLTPFTYMLEKWRMMVFKGEIPKQOMKKWEMKREIVGVBPVPHDETTC 480
Db 439 LLKQALTTVGLTPFTYMLEKWRMMVFKGEIPKQOMKKWEMKREIVGVBPVPHDETTC 498
QY 481 DPASLFHVSNDSYSPRYRTLYQFOFQALCOAAGHEGLHKCDISNSTEAGQKL 536
Db 499 DPASLFHVSNDSYSPRYRTLYQFOFQALCOAAGHEGLHKCDISNSTEAGQKL 554

;
; LENGTH: 555
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-973-1158-72

Query Match 89.7%; Score 2897; DB 6; Length 555;
Best Local Similarity 99.8%; Pred. No. 6.7e-226;
Matches 535; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

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QY      1 STIEEAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKNSAFLEKEQST 60
DB      19 STIEEAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKNSAFLEKEQST 78

QY      61 LAQMYPLOEIQNLTVKQLQALQONGSSVLSDBKSKRLNTILNTMTSTIYSTGKVCNPNP 120
DB      79 LAQMYPLOEIQNLTVKQLQALQONGSSVLSDBKSKRLNTILNTMTSTIYSTGKVCNPNP 138

QY      121 QECILLEPGLNEIMANSLDYNERLWAWESWRSEVGVKQLRPLYEYVVLKNEMARANHYED 180
DB      139 QECILLEPGLNEIMANSLDYNERLWAWESWRSEVGVKQLRPLYEYVVLKNEMARANHYED 198

QY      181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTEETFEIKPLYEHLHAYVRKLMNAYPSYISP 240
DB      199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTEETFEIKPLYEHLHAYVRKLMNAYPSYISP 258

QY      241 IGCLPAHLGDMWGRFWTNLYSLTVPPFGQKPNIDVTDAMVDAQDAQRIKFAEAKFFVSV 300
DB      259 IGCLPAHLGDMWGRFWTNLYSLTVPPFGQKPNIDVTDAMVDAQDAQRIKFAEAKFFVSV 318

QY      301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVMTDDFLTAAHHEMGH 360
DB      319 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVMTDDFLTAAHHEMGH 378

QY      361 IQDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
DB      379 IQDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438

QY      421 LKQALTIIVGTLPFTYMLEKWRWVFKGBIPKQDMKKWEMKREIVGVVPEVPDHETVC 480
DB      439 LKQALTIIVGTLPFTYMLEKWRWVFKGBIPKQDMKKWEMKREIVGVVPEVPDHETVC 498

QY      481 DPASLPHVSNDSYFIRYTRTYIQFOFQALCOAAKHEGLPKHCDISNSTEAGOKL 536
DB      499 DPASLPHVSNDSYFIRYTRTYIQFOFQALCOAAKHEGLPKHCDISNSTEAGOKL 554
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RESULT 6

US-10-518-599-24
; Sequence 24, Application US/10518599
; Publication No. US20050251873A1
; GENERAL INFORMATION:
; APPLICANT: PENNINGER, JOSEPH M.
; APPLICANT: CRACKOWER, MICHAEL A.
; TITLE OF INVENTION: ACE2 ACTIVATION FOR TREATMENT OF HEART, LUNG AND
; TITLE OF INVENTION: KIDNEY DISEASE AND HYPERTENSION
; FILE REFERENCE: SONN:064US
; CURRENT APPLICATION NUMBER: US/10/518,599
; PRIOR FILING DATE: 2004-12-17
; PRIOR APPLICATION NUMBER: PCT/CA03/00882
; PRIOR FILING DATE: 2003-06-19
; PRIOR APPLICATION NUMBER: US 60/389,709
; PRIOR FILING DATE: 2002-06-19
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 24
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Rattus rattus
US-10-518-599-24

Query Match 85.3%; Score 2757; DB 6; Length 805;
Best Local Similarity 84.2%; Pred. No. 2.2e-214;
Matches 501; Conservative 42; Mismatches 52; Indels 0; Gaps 0;

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QY      1 STIEEAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKNSAFLEKEQST 60
DB      19 SLIEEAKSFLNKNQEAEDLSYQSSLASWNYNTNITEENAKQKNEAAKWSAFYEEOQK 78

QY      61 LAQMYPLOEIQNLTVKQLQALQONGSSVLSDBKSKRLNTILNTMTSTIYSTGKVCNPNP 120
DB      79 LAQNPQLQEIQNAVITKRLKALQOGSSALSPPDKNKLQNTILNTMTSTIYSTGKVCNSMNP 138

QY      121 QECILLEPGLNEIMANSLDYNERLWAWESWRSEVGVKQLRPLYEYVVLKNEMARANHYED 180
DB      139 QECILLEPGLNEIMATSTDYNNRLWAWEGWAEBVGVKQLRPLYEYVVLKNEMARANNYED 198

QY      181 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTEETFEIKPLYEHLHAYVRKLMNAYPSYISP 240
DB      199 YGDYWRGDEYVNGVDGYDSRGQLIEDVHTTEETFEIKPLYEHLHAYVRKLMNAYPSYISP 258

QY      241 IGCLPAHLGDMWGRFWTNLYSLTVPPFGQKPNIDVTDAMVDAQDAQRIKFAEAKFFVSV 300
DB      259 IGCLPAHLGDMWGRFWTNLYSLTVPPFGQKPNIDVTDAMVDAQDAQRIKFAEAKFFVSV 318

QY      301 GLPNMTQGFWNSMLTDPGNVQKAVCHPTAWDLGKGFRIILMCTKVMTDDFLTAAHHEMGH 360
DB      319 GLPQMTPGFWTNSMLTEPGDDRVKVCPTAWDLGKGFRIILMCTKVMTDDFLTAAHHEMGH 378

QY      361 IQDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 420
DB      379 IQDMAYAAQPFLLRNGANEGFHEAVGEIMSLSAATPKHLKSIIGLLSPDFQEDNETEINF 438

QY      421 LKQALTIIVGTLPFTYMLEKWRWVFKGBIPKQDMKKWEMKREIVGVVPEVPDHETVC 480
DB      439 LKQALTIIVGTLPFTYMLEKWRWVFKGBIPKQDMKKWEMKREIVGVVPEVPDHETVC 498

QY      481 DPASLPHVSNDSYFIRYTRTYIQFOFQALCOAAKHEGLPKHCDISNSTEAGOKL 540
DB      499 DPASLPHVSNDSYFIRYTRTYIQFOFQALCOAAKHEGLPKHCDISNSTEAGOKL 558

QY      541 RLKSEPTLALENVVGAKNMVRPLNYPFLPTLTKDQNKSNFVGVGWSMTDWSY 595
DB      559 SLGNSGPWTLALENVVGRNDVFKPLNYPFLFVWLKEQNRNSTVGVGWSMTDWSY 613
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RESULT 7

US-10-518-599-4
; Sequence 4, Application US/10518599
; Publication No. US20050251873A1
; GENERAL INFORMATION:
; APPLICANT: PENNINGER, JOSEPH M.
; APPLICANT: CRACKOWER, MICHAEL A.
; TITLE OF INVENTION: ACE2 ACTIVATION FOR TREATMENT OF HEART, LUNG AND
; TITLE OF INVENTION: KIDNEY DISEASE AND HYPERTENSION
; FILE REFERENCE: SONN:064US
; CURRENT APPLICATION NUMBER: US/10/518,599
; CURRENT FILING DATE: 2004-12-17
; PRIOR APPLICATION NUMBER: PCT/CA03/00882
; PRIOR FILING DATE: 2003-06-19
; PRIOR APPLICATION NUMBER: US 60/389,709
; PRIOR FILING DATE: 2002-06-19
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 4
; LENGTH: 805
; TYPE: PRT
; ORGANISM: Mus musculus
US-10-518-599-4

Query Match 85.3%; Score 2755; DB 6; Length 805;
Best Local Similarity 84.2%; Pred. No. 3.2e-214;
Matches 501; Conservative 37; Mismatches 57; Indels 0; Gaps 0;

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QY      1 STIEEAKTFLDKFNHEADLFYQSSLASWNYNTNITEENVQNMNAGDKNSAFLEKEQST 60
DB      19 SLTEENAKTFLNPNQEAEDLSYQSSLASWNYNTNITEENAKQKNEAAKWSAFYEEOQK 78
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